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EXAMINER
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**BEFORE THE BOARD OF PATENT APPEALS
AND INTERFERENCES**

Paper No. 14

Application Number: 09/546,143

Filing Date: April 10, 2000

Appellant(s): MATZINGER ET AL.

Aaron F. Dubberley

For Appellant

EXAMINER'S ANSWER

This is in response to the appeal brief filed 10-12-01.

(1) Real Party in Interest

A statement identifying the real party in interest is contained in the brief.

(2) Related Appeals and Interferences

A statement identifying the related appeals and interferences which will directly affect or be directly affected by or have a bearing on the decision in the pending appeal is contained in the brief.

(3) Status of Claims

The statement of the status of the claims contained in the brief is correct.

(4) Status of Amendments After Final

The appellant's statement of the status of amendments after final rejection contained in the brief is correct.

(5) Summary of Invention

The summary of invention contained in the brief is correct.

(6) Issues

The appellant's statement of the issues in the brief is correct.

(7) Grouping of Claims

Appellant's brief includes a statement that claims 10-27 do not stand or fall together and provides reasons as set forth in 37 CFR 1.192©(7) and ©(8).

(8) ClaimsAppealed

The copy of the appealed claims contained in the Appendix to the brief is correct.

(9) Prior Art of Record

5,583,222

BARBIER et. al.

12-1996

Adams, C.P. et. al. "Total Synthesis of balanol: a potent protein kinase C inhibitor of fungal origin" Jounral of Chemical Society, Perkin Transaction I, 1975, pp. 2355-2362.

Krogsgaard-Larsen P. et. al. "Inhibitors of GABA Uptake. Syntheses and H NMR Spectroscopic Investigations of Guvacine, (3RS, 4SR)-4-hydroxypiperidine-3-carboxylic Acid, and Related Compounds" Acta Chemica Scandinavica B, vol. 32, no. 5 (1978), pp. 327-334.

Wade, G. L. Jr. ORGANIC CHEMISTRY, © 1987, pages 103 and 115, Prentice Hall Inc., Englewood Cliffs, New Jersey (cited as a basic principle for geometric isomerism – not to be construed as a new prior art).

(10) First Ground of Rejection

The following ground(s) of rejection are applicable to the appealed claims:

Claim Rejections - 35 USC § 112

The following is a quotation of the **second** paragraph of 35 U.S.C. 112:

The specification shall conclude with one or more claims particularly pointing out and distinctly claiming the subject matter which the applicant regards as his invention.

Claims 12, 14, 15, 17, 19, 21, 23, and 25 are rejected under 35 U.S.C. 112, second paragraph, as being indefinite for failing to particularly point out and distinctly claim the subject matter which applicant regards as the invention. Said claims recite the limitation of "amino protecting group" which has no description in the specification other than "tert.-butoxycarbonyl" as a sole representative of said group. Thus, one skilled in the art cannot ascertain what other groups can be considered as an "amino protecting group". Therefore, the **metes and bounds** of the invention is indefinite.

The following is a quotation of the **first** paragraph of 35 U.S.C. 112:

The specification shall contain a written description of the invention, and of the manner and process of making and using it, in such full, clear, concise, and exact terms as to enable any person skilled in the art to which it pertains, or with which it is most nearly connected, to make and use the same and shall set forth the best mode contemplated by the inventor of carrying out his invention.

Claims 12, 14, 15, 17, 19, 21, 23, and 25 are rejected under 35 U.S.C. 112, first paragraph, because the specification, while being enabling for R^4 as a tert-butyl ester, tert-

butyl carboxylate, or tert-butoxycarbonyl, does not reasonably provide enablement for the genera with R^4 as another functional group serving as an “amino protecting group”. The specification does not enable any person skilled in the art to which it pertains, or with which it is most nearly connected, to make and use the invention commensurate in scope with these claims. The disclosure does not provide guidance as to what functional groups, and/or rings can be considered as an amino protecting group. Its mere statement “ R^4 is a protecting group, preferably a tert.-butoxycarbonyl group” is insufficient for one skilled in the art to consider what other groups can be an “amino protecting group”. In considering enablement, undue experimentation is an important factor. Here, the scope of “amino protecting group” is undoubtedly broad, and the generic teaching for preparation only gears toward one “amino protecting group”, namely, tert.-butoxycarbonyl. Thus, regarding other groups, one skilled in the art will have to carry out undue experimentation, as the chemical art is unpredictable.

Note, the Federal Circuit has repeatedly held that “**the specification must teach those skilled in the art how to make and use the full scope of the claimed invention without ‘undue experimentation’**” (see **In re Wright**, 999 F. 2d 1557, 1561, 27 U.S.P.Q. 2d 1510, 1513 (Fed. Cir. 1993)). Also, the disclosure does not provide the starting material for R^4 , nor a source for an “amino protecting group”, and thus, undue experimentation is inevitable for one skilled in the art to make and use compounds with R^4 as a group other than tert.-butoxycarbonyl. See **In re Howarth**, 210 U.S.P.Q 689, 693 regarding insufficient enablement. Note, in said case, the starting material was not disclosed, nor was a source named. The court, then, ruled that “**burden rests upon applicant who chooses to rely upon**

general knowledge in the art to render his disclosure enabling to establish that those of ordinary skill in the art can be expected to possess or know where to obtain this knowledge;..." Thus, 'no disclosure of starting material' is a sound reason to render a specification with insufficient enablement.

(11) Response to Argument on the above issues of 112/ 1st and 2nd paragraphs for claims 12, 14, 15, 17, 19, 21, 23, and 25

Applicant disputes the indefiniteness of the terms "amino protecting group" for the following reasons:

- a. Said terms are well-known in the art;
- b. Over 500 US patents recite said terms in their claims.
- c. Citing textbook of Green along with case laws such as: In re Skoll (187 USPQ 481), In re Fuetterer (138 USPQ 217), In re Bowen (181 USPQ 48), In re Robins (166 USPQ 552) to support applicant's position.
- d. The definition of R⁴ as an "amino protecting group" is a functional language.
- e. As for enablement, applicant feels that it is not necessary to teach what is well known in the art.

Despite applicant's argument, the issues of 112/1st and 2nd paragraphs should be upheld for the following reasons:

- i. The issue of indefiniteness is not whether one skilled in the art can understand a term (or terms), rather it is the metes and bounds of the invention. In the instant case, the terms “amino protecting group” as defined for R⁴ does not clearly define the scope of claims 12, 14, 15, 17, 19, 21, 23, and 25. The disclosure does not describe what functional groups, ring(s) said terms encompass, except tert.-butoxycarbonyl. Therefore, when interpreting the instant claims in light of the specification, one does not have a guidance as to what is claimed (or **not** claimed) by the terms “amino protecting group”. Applicant cites the textbook of Green for description of an “amino protecting group”. However, it is still unclear whether the scope of “amino protecting group” includes groups cited by Green, or goes beyond that. While breadth is not indefiniteness, the claims must particularly point out and **distinctly define the metes and bounds of the subject matter that will be protected by the patent grant.**
- ii. It is recognized that the terms “amino protecting group” are recited in a lot of US patents. However, usually, said terms are used specifically in a reaction step in a claimed process. When read in light of a process, said terms have a more definite metes and bounds. In the instant case, applicant claims intermediates (i.e., compounds) having “amino protecting group” as a substituent. Because in the specification, there is no description as to what constitutes said group (other than tert.-butoxycarbonyl), the final structure of a claimed intermediate is indeterminate. Therefore, the scope of claims reciting said terms is indefinite.

iii. Case laws cited by applicant are outdated. The most recent case law is (**Genetech Inc. v. Novo Nordisk**, 108 F.3d 1361, 42 USPQ 2d 1001 (Fed. Cir. 1997)), in which the court ruled that relying on the knowledge of one skilled in the art cannot cure the deficiency in enablement. Just because a term is well-known in the art, it does not mean one skilled in the art can prepare any intermediate having any “amino protecting group”. Based on what disclosed by applicant, one skilled in the art can only make compounds with R⁴ as tert.-butoxycarbonyl. It is true applicant does not have to provide enablement for known processes in the art, but applicant is obligated to provide **enablement in commensurate with the scope of the claims**. In the instant case, because the terms “amino protecting group” have such a broad scope that the enablement for only tert.-butoxycarbonyl does **not sufficiently** guide the skilled chemist in preparing intermediates with other amino protecting groups.

(12) Second Ground of Rejection

The following ground(s) of rejection are applicable to the appealed claims:

Claim Rejections - 35 USC § 102

The following is a quotation of the appropriate paragraphs of 35 U.S.C. 102 that form the basis for the rejections under this section made in this Office action:

A person shall be entitled to a patent unless –

(b) the invention was patented or described in a printed publication in this or a foreign country or in public use or on sale in this country, more than one year prior to the date of application for patent in the United States.

(e) the invention was described in a patent granted on an application for patent by another filed in the United States before the invention thereof by the applicant for patent, or on an international application by another who has fulfilled the requirements of paragraphs (1), (2), and (4) of section 371© of this title before the invention thereof by the applicant for patent.

The rejection of claim 23 under 35 U.S.C. 102(a) based on **Lamp et. al.** is withdrawn herein in light of applicant's argument.

102(b) – Claims 17, 23, and 25 are rejected under 35 U.S.C. 102(b) as being anticipated by the following references:

- a. **Krogsgaard-Larsen et. al.** (Acta Chem. Scan. B, 32 (1978), pp. 327-334): Compounds 12 and 13 on page 328 are embraced by formula VIII in claim 17 with R³ as lower alkyl, and R⁴ as an amino protecting group.
- b. **Adams et. al.** (J. Chem. Soc. Perkin Trans. I, 1995, pp. 2355-2362): Formula X in claim 23 inherently embraces compound 20 on page 2356. Formula XI in claim 25 inherently embraces compound 21 on page 2357.

102(e) – Claim 25 is rejected under 35 U.S.C. 102(e) as being anticipated by **Barbier et. al.** (US 5,583,222). Formula XI in claim 25 inherently embraces compounds B1-B23 listed on columns 18-21.

(13) Response to Argument on 102(b), and (e) rejections for claims 17, 23, and 25

Applicant disputes that the 102 rejections are improper because said references do not disclose the cis-form of the claimed formulae. Applicant argues that the trans-form is structurally different from the cis-form, and thus, cannot anticipate the cis-form of the claimed formulae.

Applicant's argument is well taken; however, the 102 rejections based on inherency should be upheld for the following reasons:

- i. Despite applicant's assertion, cis- and trans- forms of a compound have **similar structures** due to their common cores, and substituents. They only differ in the spatial orientation of their substituents. However, spatial orientation of a compound can flip flop from one form to the other because **bonds are not static**. Anyone skilled in the art would know that if a trans-form of a compound, a cis-form also **exists inevitably**. This is the most fundamental principle in stereo-chemistry. So, if a reference discloses a trans-form, then a cis-form will be inherently embraced.
- ii. Regarding claim 17, compound #12 of Krogsgaard-Larsen clearly a **cis**-compound because both the –OH locates on the same side with –C(O)OCH₃ just as the claimed formula VIII. It is not a racemic mixture of cis- and trans- as suggested by applicant. The symbol, “(+)”, refers to optical isomers of the **cis**-compound. Note, both the –OH and the –C(O)OCH₃ can be pointed upward or downward, and still have **cis**-

configuration. See Wade's ORGANIC CHEMISTRY, pages 103 and 115, as evidence for the basic principle of cis- and trans- configurations (or geometric isomerism).

(14) Third Ground of Rejection

The following ground(s) of rejection are applicable to the appealed claims:

Claim Rejections - 35 USC § 103

The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:

(a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negated by the manner in which the invention was made.

Claims 25 and 26 are rejected under 35 U.S.C. 103(a) as being unpatentable over **Barbier et. al. (US 5,583,222)**. On column 9, Barbier et. al. disclose a group of intermediates represented by formula III which resembles the claimed formula XI since the reference's variable Y can be taken as -NH while the -CO-A corresponds to the instant variable R¹, and R¹⁶ (represents an "amino protecting group") corresponds to the instant R⁴. Furthermore, column 18 lists intermediates B1-B23 which have tert-butyl carboxylate as an amino protecting group (e.g., see intermediate B4). While Barbier et. al. do not disclose the cis-configuration of compounds of formula III or its species, such form is suggested in the racemic mixture of cis- and trans- represented by formula III.

Furthermore, Barbier et. al. recognizes that substituents on the heterocyclic ring can have cis-configuration as well (see column 4, line 30). Therefore, one of the ordinary skill in the art would have been motivated to make the cis-configuration of compounds of formula III because such a configuration had been acknowledged by Barbier et. al. as an alternative to trans-configuration. Thus, at the time of the invention, it would have been obvious to one skilled in the art to make intermediates of formula XI and its species because one skilled in the art can always resolve a racemic mixture of formula III by conventional methods to obtain the cis-form of formula XI claimed herein.

(15) Response to Argument on the 103 rejection for claims 25 and 26

Applicant's argues that the broad genus of the disclosed formula III does not allow one skilled in the art to select the intermediates claimed herein.

Applicant's argument cannot overcome the rejection for the following reasons:

- i. Barbier et. al. not only disclose the genus of formula III, but also its species (B1-B23).
So, one cannot dispute the teaching of Barbier et. al. as merely being broad.
- ii. Because the disclosed species B1-B23 are in trans-form, formula III was cited in the rejection as a racemic mixture, which suggests that a cis-form is also implicitly suggested. Therefore, it is within the level of one skilled in the art to obtain the claimed cis-form from the teaching of Barbier et. al., and conventional methods of resolving cis- and trans- forms.

Art Unit: 1624

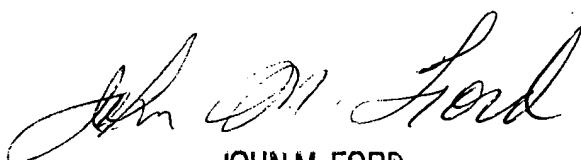
For the above reasons, it is believed that all rejections should be sustained.

Respectfully submitted,

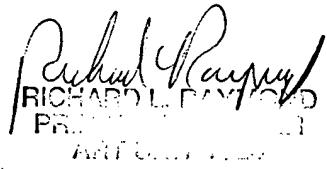


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March 27, 2003



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Total synthesis of balanol: a potent protein kinase C inhibitor of fungal origin

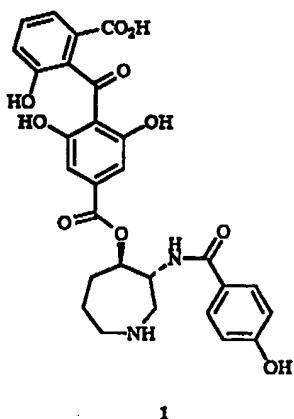
C. P. Adams,^a S. M. Fairway,^a C. J. Hardy,^a D. E. Hibbs,^b M. B. Hursthouse,^b A. D. Morley,^a B. W. Sharp,^a N. Vicker^{*,a} and I. Warner^a

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The total synthesis of the fungal metabolite balanol, a potent inhibitor of protein kinase C, is described. The synthesis includes a novel synthesis of 3-amino-4-hydroxyazepanes via a directed ring expansion of 3-bromopiperidin-4-ones.

Banol **1** is a structurally novel inhibitor of protein kinase C, a family of phospholipid dependent serine/threonine protein kinases which play an important role in cell growth, signal transduction and differentiation.¹ The activated enzyme² has been implicated in many diseases, such as cancer, inflammation and HIV infection; therefore inhibitors of protein kinase C may be therapeutic.³ Balanol **1** was initially isolated by workers at Sphinx Pharmaceuticals from *Verticillium balanoides*^{4,5} and more recently from species of *Fusarium*⁶ by a team at Nippon Roche. Synthetic interest in balanol is intense, owing to its chemical structure, biological activity and its low availability from natural sources. As our synthetic route was nearing completion, the total synthesis of balanol was recently reported by workers at Sphinx,⁷ and by the Nicolaou group.⁸ Here we describe a new synthesis of balanol with novel routes to the hexahydroazepine (azepane) and benzophenone portions.



Results and discussion

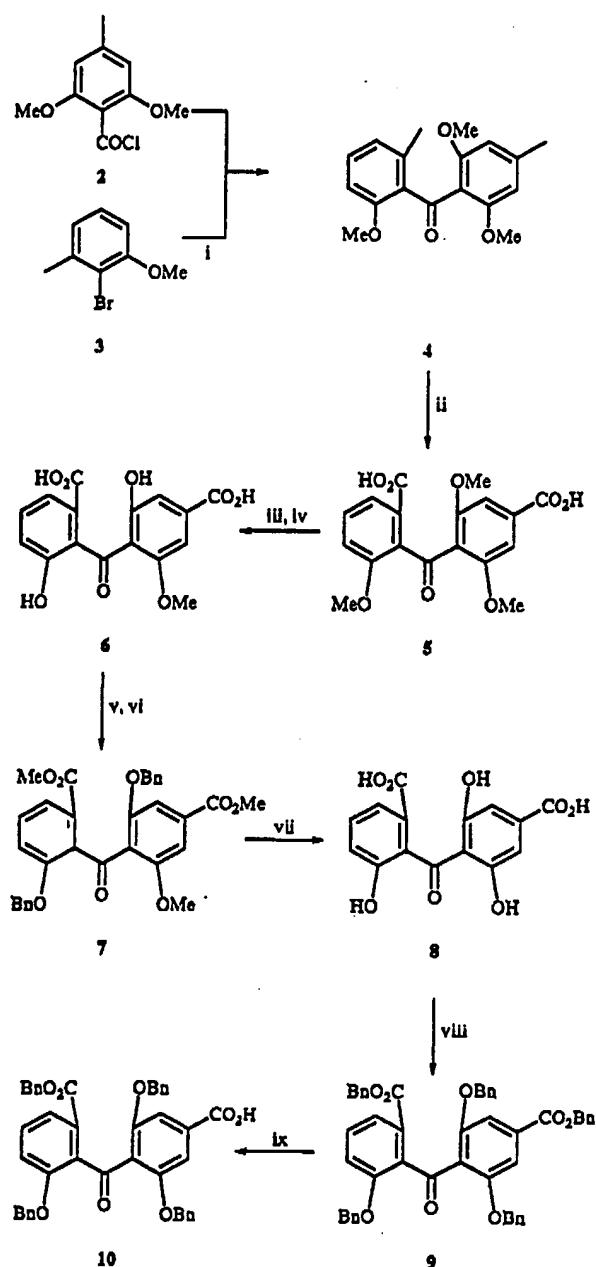
The key disconnection for the synthesis of balanol is the ester bond joining the azepane and the benzoic acid. This implies that preparation of suitably protected benzophenone and azepane portions, which would be coupled together to form the ester bond, could then be deprotected to give balanol **1**.

The synthesis of the benzophenone portion started from 2,6-dimethoxy-4-methylbenzoyl chloride⁹ **2** and 2-bromo-3-methoxytoluene¹⁰ **3** which were readily prepared from commercially available 3,5-dimethoxytoluene and 2-amino-3-methoxytoluene respectively, in 86 and 85% yield. Formation of the Grignard reagent of **3** and reaction of this with **2** gave the hindered

benzophenone **4** in 80% yield. Oxidation of the methyl groups of **4** using potassium permanganate gave the diacid **5** in 38% yield. The diacid **5** had limited solubility and was quantitatively converted into the dimethyl ester. Treatment of this dimethyl ester with boron tribromide gave **6** in 95% yield with only small amounts of **8** being detected. Further demethylation of **6** to give the triphenolic diacid **8** directly was problematic and treatment of **6** with boron tribromide and a host of other demethylating agents yielded only traces of **8**. Possibly once **6** has been formed, its insolubility and ability to complex with reagents hinders the final demethylation. This problem was overcome by re-esterification of **6** to give the dimethyl ester followed by benzylation of the phenolic groups to give **7** in 74% overall yield. Treatment of **7** with boron tribromide in dichloromethane gave the desired triphenolic diacid **8** in 52% yield and **6** in 41% yield which could be recycled. In this case, the demethylation occurred competitively with the more hindered benzyl groups, which, since they are more labile than the methoxy group, are then subsequently removed. Complete demethylation of the dimethyl ester of **6** under the above conditions was unsuccessful, only ester hydrolysis being observed. The triphenolic diacid **8** was converted into the pentabenzyl compound **9** in 42% yield selective hydrolysis of which gave **10** in 91% yield. This compound was identical spectroscopically with an authentic sample prepared by a different route.^{7,8} The tetrabenzyl compound **10** is suitably protected for activation and coupling to a protected portion of the azepane **21** to give balanol.

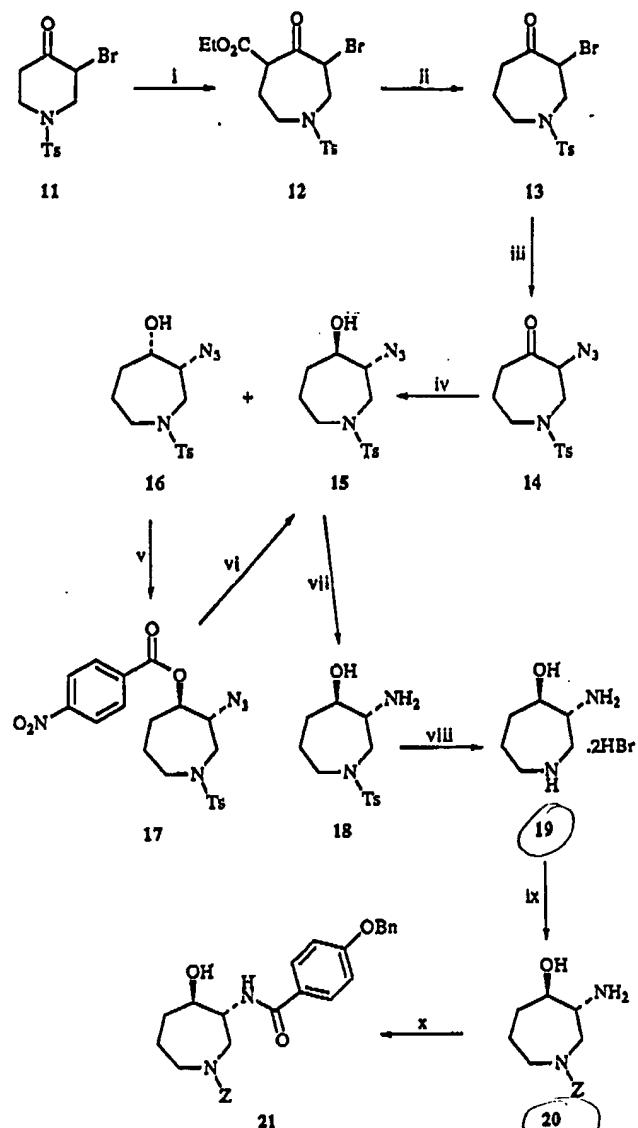
The synthesis of the azepane portion of balanol started from commercially available piperidin-4-one. *N*-Tosylation under standard conditions followed by bromination gave **11** in 80% overall yield. The tosyl group was chosen to protect the nitrogen as this group was more stable to the bromination conditions and subsequent stages than carbamates and all compounds containing the tosyl group were crystalline. It was expected that the tosyl group would have to be removed and replaced by a more labile group after the requisite functional group manipulations had been completed so that deprotection at the final stage of the balanol synthesis would not be problematic.

The regiospecific homologation of unhindered α -bromo ketones using ethyl diazoacetate and boron trifluoride-diethyl ether has been reported.¹¹ The observed regioselectivity of homologation has been demonstrated to be due to a combination of steric and electronic factors.¹¹ The electron-withdrawing effect of the α -bromo substituent suppresses the migration of the carbon atom bearing it and the steric effect of the bromine atom enhances this selectivity of migration. This



Scheme 1 *Reagents and conditions:* i, Mg, THF, 2 (80%); ii, KMnO₄, pyridine (aq.) (38%); iii, SOCl₂, MeOH, iv, Br₂, CH₂Cl₂ (95% from 5); v, SOCl₂, MeOH; vi, NaH, BnBr, DMF (74% from 6); vii, BBr, CH₂Cl₂ (52%); viii, NaH, BnBr, DMF (42%); ix, Na₂CO₃ (aq.), EtOH (91%)

sort of regiospecific homologation has not been applied previously to unsymmetrical ketones present in heterocyclic systems. A novel use of this type of ring expansion was demonstrated in the conversion of 11 into 12 using ethyl diazoacetate under Lewis acid conditions, which proceeded in 71% yield. As predicted, insertion of the methylene was selective for the unhindered side of the ketone, with none of the other regioisomer being observed. Hydrolysis and decarboxylation of 12 gave 13 in 90% yield. Reaction of 13 with sodium azide resulted in nucleophilic displacement of bromide by azide to give 14 in 73% yield. Reduction of 14 with sodium boranide (NaBH₄) gave a *cis:trans* (2.4:1) mixture of azido alcohols 16 and 15 which were easily separable by chromatography giving 15 in 26% and 16 in 62% yields, respectively. Conversion of the major unwanted *cis* isomer 16 into the *trans* isomer 15



Scheme 2 *Reagents and conditions:* i, EtO₂C-N₃, Br, CH₂Cl₂ (71%); ii, HCl (aq.), dioxane (90%); iii, Na₃N, AcOH, DMF (73%); iv, NaBH₄, EtOH (88%); v, PPh₃, DIAD, THF, *p*-nitrobenzoic acid (84%); vi, NaOH (aq.), MeOH, dioxane (99%); vii, LiAlH₄, THF (85%); viii, HBr (aq.) (68%); ix, Et₃N, CH₂Cl₂, 18-crown-6, benzyl chloroformate (88%); x, 4-(benzyloxy)benzoyl chloride, Et₃N, CH₂Cl₂ (65%)

proceeded in 84% yield using a Mitsunobu¹² inversion *via* the *p*-nitrobenzoate 17 followed by hydrolysis. The *trans* relationship of the hydroxy and azido groups in 15 was unambiguously established by X-ray crystallography at this stage as shown in Fig. 1. Treatment of 15 with lithium aluminium hydride gave the 3-amino-4-hydroxyazepane 18 in 85% yield. At this stage, the *N*-tosyl group had to be replaced by a group that would be more easily removed in the final stage of the synthesis. Removal of the *N*-tosyl group was effected in 68% yield using aqueous hydrobromic acid to give 19. Selective protection of the secondary amino group in 19 proceeded in 88% yield using benzyl chloroformate in the presence of 18-crown-6 to give 20. Reaction of 20 under standard conditions with 4-(benzyloxy)benzoyl chloride gave 21 in 65% yield, which is a suitably protected azepane portion of balanol to be coupled to 10 following activation. The enantiomers of 21 were resolved by forming their Mosher's esters¹³ 22 followed by chromatography and hydrolysis to give both enantiomers of 21 in

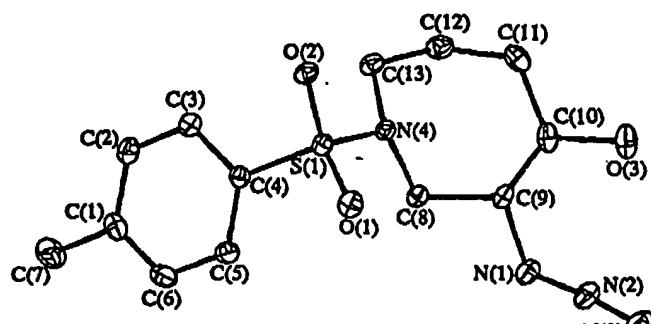
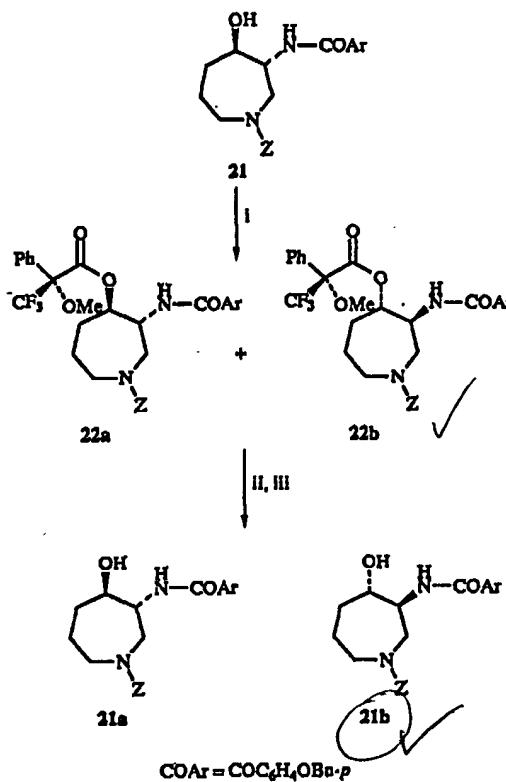
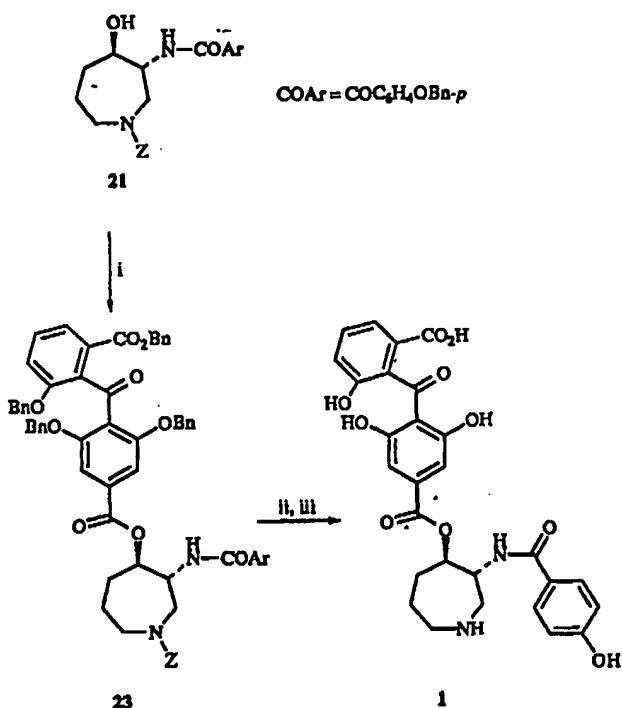


Fig. 1 X-Ray structure of compound 15

Scheme 3 Reagents and conditions: i, acid chloride of (S)-(-)-MTPA; CH_2Cl_2 , Et_3N , DMAP; ii, chromatography (96% from 21); iii, KOH , MeOH (100%)

96% overall yield and in >99% ee. The enantiomeric purity was checked relative to the racemate using chiral chromatography. This separation allowed access to both enantiomerically pure forms of balanol in a similar manner to that previously described.⁷ Compound 21 was identical with an authentic sample prepared by a different route.⁸

Activation of 10 by the method of Mukaiyama,¹⁴ as employed in the Nicolaou synthesis⁸ of balanol followed by coupling to 21 gave 23 which was identical spectroscopically with an authentic sample.⁸ Hydrogenolysis of 23, with palladium black in aqueous acetic acid-ethyl acetate gave balanol 1 as a major product. In our hands the use of THF as a co-solvent under similar conditions to Nicolaou during the deprotection step gave rise to quantities of *N*-hydroxybutylbalanol. This side reaction was avoided by replacing the THF with ethyl acetate. Purification of 1 using reverse phase HPLC gave pure balanol 1 identical by HPLC, MS and NMR with an authentic sample of balanol.⁸

Scheme 4 Reagents and conditions: i, 10, 2-chloro-1-methylpyridinium iodide, DMAP, Et_3N , CH_2Cl_2 (37%); ii, H_2 , Pd black, HOAc , EtOAc , H_2O ; iii, HPLC (53% from 23)

Experimental

Mps were determined using an Electrothermal apparatus and are uncorrected. IR spectra were recorded on a Nicolet 20SXB spectrophotometer. ^1H NMR spectra were obtained using a Varian VXR400 instrument (400 MHz), δ values quoted are relative to internal TMS and J values are given in Hz. Mass spectra were measured with either a Varian VG 7070E or Finnegan TSQ 700 spectrometer. Flash chromatography was performed using Sorbsil C 60 (40–60 μm mesh) silica gel. Analytical thin layer chromatography was carried out on 0.25 mm precoated silica gel plates (E. Merck Kieselgel 60 F₂₅₄) and compounds were visualised using UV fluorescence, ethanolic phosphomolybdc acid or aqueous potassium permanganate.

2,6-Dimethoxy-4-methylphenyl 2-methoxy-6-methylphenyl ketone 4

A suspension of magnesium turnings (0.6 g, 25 mmol) in THF (2 cm^3) under an atmosphere of nitrogen was treated with 1,2-dibromoethane (0.14 cm^3). The mixture was slowly stirred and heated to reflux whilst a solution of 2-bromo-3-methoxytoluene 3 (5.03 g, 25 mmol) in THF (20 cm^3) was added dropwise. The mixture was heated at reflux for 1 h, allowed to cool to room temperature and then filtered to remove any traces of magnesium. To this Grignard reagent a solution of 2,6-dimethoxy-4-methylbenzoyl chloride 3 (5.25 g, 25 mmol) in THF (20 cm^3) was added dropwise, the temperature of the mixture being kept at *ca.* 25 °C. The mixture was stirred at room temperature for 30 min after which the solvent was removed under reduced pressure to give a pale red syrup which was treated with saturated aq. NH_4Cl (200 cm^3) and ethyl acetate (150 cm^3). The layers were separated and the aqueous layer was extracted with ethyl acetate (50 cm^3). The combined organic layer and extracts were washed with 5% aq. NaHCO_3 (3 \times 100 cm^3) and brine (100 cm^3), dried (MgSO_4), filtered and evaporated under reduced pressure to yield a yellow solid (7.2

g). This was recrystallised from ethanol to yield 4 (6.0 g, 80%) as a white solid, mp 132–133 °C (Found: C, 71.8; H, 6.6. $C_{18}H_{20}O_4$ requires C, 72.0; H, 6.7%); ν_{max} (KBr)/cm⁻¹ 1670s (CO); δ_H (CDCl₃) 2.31 (3 H, s, Me), 2.34 (3 H, s, Me), 3.58 (3 H, s, 6'-OMe), 3.66 (6 H, s, 2'- and 6'-OMe), 6.34 (2 H, s, 3'- and 5'-H), 6.67 (1 H, d, J 8), 6.78 (1 H, d, J 8) and 7.16 (1 H, t, J 8, 4-H); *m/z* 301 (MH⁺).

4-Carboxy-2,6-dimethoxyphenyl 2-carboxy-6-methoxyphenyl ketone 5

To a stirred solution of potassium permanganate (28 g, 3 equiv.) in water (160 cm³) and pyridine (160 cm³) at *ca.* 100 °C, aq. NaOH (20%; 4 cm³) was added. To this stirred mixture a solution of 4 (17.67 g, 16.98 mmol) in pyridine (90 cm³) was added dropwise followed by water (90 cm³). The mixture was heated at reflux for 1 h after which further quantities of potassium permanganate (28 g, 3 equiv.) were added hourly to it, stirring and heating being continued, until a total of 208 g, 24 equiv. had been added. The hot reaction mixture was filtered through Celite and the filter cake was washed with water (4 × 150 cm³) and pyridine (50 cm³). The filtrate was cooled on ice and acidified to pH 1 by the addition of concentrated aq. HCl; any precipitated solid was filtered off. The acidic aqueous filtrate was treated with NaCl (200 g) and extracted with ethyl acetate (6 × 400 cm³). The combined extracts were dried (MgSO₄), filtered and evaporated under reduced pressure and the residue treated with diethyl ether to give 5 (8.1 g, 38%) as a white solid, mp 253–256 °C (Found: C, 59.1; H, 4.5. $C_{18}H_{16}O_8 \cdot 0.25H_2O$ requires C, 59.25; H, 4.4%); ν_{max} (KBr)/cm⁻¹ 3400br and 2900br (OH), 1690s (CO); δ_H (CD₃SOCD₃) 3.59 (3 H, s, 6'-OMe), 3.64 (6 H, s, 2'- and 6'-OMe), 7.17 (2 H, s, 3'- and 5'-H), 7.22 (1 H, d, J 8), 7.29 (1 H, d, J 8) and 7.46 (1 H, t, J 8, 4-H); *m/z* 361 (MH⁺).

4-Carboxy-2-hydroxy-6-methoxyphenyl 2-carboxy-6-hydroxyphenyl ketone 6

To a stirred solution of 5 (1.01 g, 2.8 mmol) in methanol (25 cm³), thionyl chloride (1.17 cm³, 16 mmol) was added dropwise and the mixture heated at reflux for 2 h. The mixture was allowed to cool to room temperature and then evaporated to dryness. Treatment of the residue with toluene (2 × 20 cm³) and removal of the solvent gave a white solid (1.09 g) which was dissolved in dichloromethane (37 cm³) and the resulting solution was cooled to –60 °C. To the solution at –60 °C boron tribromide in dichloromethane (1 mol dm⁻³; 31 cm³) was added dropwise and the mixture was allowed to warm to room temperature with stirring for 18 h. The mixture was cooled to 0 °C and water (50 cm³) was added dropwise to it. The mixture was allowed to warm to room temperature and any precipitated solid was removed by filtration. The layers were separated and the aqueous layer was adjusted to pH 4 by the addition of aq. NaHCO₃ (5%) and the aqueous layer was re-extracted with ethyl acetate (2 × 50 cm³). The solid removed by the previous filtration was dissolved in the combined organic extracts and the solution dried (MgSO₄), filtered and evaporated under reduced pressure to yield 6 (0.9 g, 95%) as a pale yellow solid, mp 262–265 °C (Found C, 57.8; H, 3.7. $C_{16}H_{12}O_8$ requires C, 57.8; H, 3.6%); ν_{max} (KBr)/cm⁻¹ 3490br, 3415br and 2900br (OH), 1680s and 1635s (CO); δ_H (CD₃SOCD₃) 3.49 (3 H, s, 6'-OMe), 6.90 (1 H, s), 7.05 (1 H, s), 7.08 (1 H, d, J 8), 7.32 (1 H, t, J 8, 4-H), 7.39 (1 H, d, J 8), 9.90 (1 H, br s, OH) and 12.75 (1 H, br s; OH); *m/z* 333 (MH⁺).

2-Hydroxy-6-methoxycarbonylphenyl 2-hydroxy-6-methoxy-4-methoxycarbonylphenyl ketone

To a stirred solution of 6 (5.64 g, 17 mmol) in methanol (200 cm³) thionyl chloride (12 cm³, 165 mmol) was added dropwise. The mixture was heated at reflux for 3 h, after which it was

allowed to cool to room temperature and then evaporated to dryness to give a light brown crystalline solid. This was then treated first with toluene (2 × 20 cm³) which was evaporated under reduced pressure and then with diethyl ether to give a cream crystalline solid (6.13 g, 100%), mp 154–156 °C (Found C, 59.8; H, 4.5. $C_{16}H_{16}O_8$ requires C, 60.0; H, 4.5%); ν_{max} (KBr)/cm⁻¹ 3450br (OH), 1725s and 1710s (CO); δ_H (CDCl₃) 3.46 (3 H, s, 6'-OMe), 3.52 (3 H, s, CO₂Me), 3.83 (3 H, s, CO₂Me), 6.94 (1 H, s), 7.12 (1 H, d, J 8), 7.30 (1 H, s), 7.38 (1 H, t, J 8, 4-H), 7.44 (1 H, d, J 8); *m/z* 361 (MH⁺).

2-Benzylxyloxy-6-methoxycarbonylphenyl 2-benzylxyloxy-6-methoxy-4-methoxycarbonylphenyl ketone 7

To a stirred solution of 2-hydroxy-6-methoxycarbonylphenyl 2-hydroxy-6-methoxy-4-methoxycarbonylphenyl ketone (6.13 g, 17 mmol) in dry DMF (120 cm³) at 0 °C NaH (60% in oil; 2.04 g, 51 mmol) was added during 3 min. After the mixture had been allowed to warm to room temperature with stirring over 1 h benzyl bromide (6.5 cm³, 51 mmol) was added to it and the reaction mixture was heated at 65 °C for 2 h. After the mixture had been cooled to 0 °C it was treated with methanol (10 cm³) and then evaporated under reduced pressure; the residue was then treated with ice water (400 cm³) and extracted with diethyl ether (2 × 150 cm³). The combined extracts were dried (MgSO₄) filtered and evaporated under reduced pressure to yield a yellow solid. Treatment of this with diethyl ether gave 7 (7 g, 74%) as a cream solid, mp 155–158 °C (Found C, 71.1; H, 5.1. $C_{32}H_{28}O_8$ requires C, 71.1; H, 5.2%); ν_{max} (KBr)/cm⁻¹ 1730s and 1670s (CO); δ_H (CDCl₃) 3.64 (3 H, s, 6'-OMe), 3.70 (3 H, s, CO₂Me), 3.95 (3 H, s, CO₂Me), 4.75 (2 H, s, OCH₂Ph), 4.86 (2 H, s, OCH₂Ph), 6.94 (3 H, m), 7.05 (2 H, m), 7.10 (1 H, s) and 7.15–7.35 (9 H, m); *m/z* 541 (MH⁺).

4-Carboxy-2,6-dihydroxyphenyl 2-carboxy-6-hydroxyphenyl ketone 8

To a stirred solution of 7 (6 g, 11.1 mmol) in dichloromethane (150 cm³) at –60 °C boron tribromide in dichloromethane (1 mol dm⁻³; 122 cm³) was added dropwise during 45 min. The reaction mixture was allowed to warm to room temperature after which it was kept at ambient temperature for 2 days. It was then cooled to 10 °C, treated dropwise with water (150 cm³) and stirred at room temperature for 1 h. The layers were separated and the aqueous layer adjusted to pH 4 by the addition of solid NaHCO₃. The aqueous layer was extracted with ethyl acetate (2 × 100 cm³) and the combined extracts were dried (MgSO₄), filtered and evaporated under reduced pressure to yield a yellow solid. The residue was purified by column chromatography on silica, eluting with methanol–dichloromethane (5:95, v/v), to give 8 (1.85 g, 52% uncorrected) as a light yellow solid, mp 222–225 °C (Found C, 56.3; H, 3.2. $C_{15}H_{10}O_8$ requires C, 56.6; H, 3.2%); ν_{max} (KBr)/cm⁻¹ 3500br, 3100br, 2900br and 2600br (OH), 1700s and 1640s (CO); δ_H (CD₃SOCD₃) 6.69 (2 H, s, 2'- and 6'-H), 7.06 (1 H, d, J 8), 7.27 (1 H, t, J 8, 4-H) and 7.48 (1 H, d, J 8; *m/z* 318 (M⁺). A quantity of 6 was also recovered (1.5 g, 41%) which could be recycled.

2-Benzylxyloxy-6-benzylxyloxy-2,6-dibenzylxyloxy-4-methoxycarbonylphenyl ketone 9

To a stirred solution of 8 (0.4 g, 1.26 mmol) in dry DMF (30 cm³) at 0 °C NaH (60% in oil; 0.3 g, 7.56 mmol) was added portionwise and the mixture was allowed to warm to room temperature with stirring for 1 h. After treatment with benzyl bromide (0.9 cm³, 7.57 mmol), the mixture was heated at 65 °C for 20 h. The mixture was cooled to 0 °C and treated with a further quantity of NaH (0.15 g, 3.78 mmol) after which it was stirred at room temperature for 30 min, and then treated with benzyl bromide (0.9 cm³, 7.57 mmol). After being heated at 65 °C for a further 4 h, the mixture was cooled to 0 °C and

treated with methanol (10 cm³); it was then evaporated under reduced pressure. The residue was treated with water (60 cm³) and then extracted with ethyl acetate (3 × 50 cm³). The combined extracts were dried (MgSO₄), filtered and evaporated under reduced pressure. The residue was chromatographed on a Dynamax silica column (20 mm × 300 mm) eluting with heptane–ethyl acetate (85:15, v/v) to yield 9 (0.4 g, 42%) as a white solid, mp 127–130 °C (Found C, 77.9; H, 5.1. C₃₀H₄₀O₈ requires C, 78.1; H, 5.2%); $\nu_{\text{max}}(\text{KBr})/\text{cm}^{-1}$ 1720s and 1680s (CO); $\delta_{\text{H}}(\text{CDCl}_3)$ 4.70 (2 H, s, 6-OCH₂Ph), 4.78 (4 H, s, 2'- and 6'-OCH₂Ph), 5.12 (2 H, s, 2-CO₂CH₂Ph), 5.38 (2 H, s, 4'-CO₂CH₂Ph), 6.82 (2 H, m), 6.95 (1 H, d, J 8) and 7.01–7.48 (27 H, m); m/z 769 (MH⁺).

2-Benzoyloxy-6-benzoyloxycarbonylphenyl 2,6-dibenzoyloxy-4-carboxyphenyl ketone 10

To a stirred mixture of 9 (0.4 g, 0.52 mmol) in ethanol (30 cm³), a solution of Na₂CO₃ (0.24 g, 1.04 mmol) in water (30 cm³) was added and the mixture heated at reflux for 4 h. The mixture was allowed to cool to room temperature after which the ethanol was removed under reduced pressure. Water (150 cm³) was added to the residue and the solution adjusted to pH 1 by the addition of concentrated aq. HCl. The aqueous mixture was then extracted with ethyl acetate (3 × 50 cm³) and the combined extracts were dried (MgSO₄), filtered and the solvent was evaporated under reduced pressure to yield a pale yellow solid. Trituration of this with diethyl ether gave 10 (320 mg, 91%) as a white solid, mp 132–134 °C; $\delta_{\text{H}}(\text{CDCl}_3)$ 4.72 (2 H, s, 6-OCH₂Ph), 4.80 (4 H, s, 2'- and 6'-OCH₂Ph), 5.14 (2 H, s, 2'-CO₂CH₂Ph), 6.85 (2 H, d, J 8), 6.97 (1 H, d, J 8), 7.05–7.08 (4 H, m), 7.13 (2 H, t, J 8, 4-H) and 7.18–7.34 (16 H, m); m/z 679 (MH⁺). Spectroscopic data of 10 were identical with those reported.^{7,8}

3-Bromo-1-(4'-methylphenylsulfonyl)piperidin-4-one 11

To a solution of 1-(4'-methylphenylsulfonyl)piperidin-4-one (262.2 g, 1.036 mol) in dichloromethane (6.5 dm³) at –5 °C, a solution of bromine (51.8 cm³, 1.004 mol) in dichloromethane (1 dm³) was added dropwise during 2 h. The temperature of the mixture was maintained between –4 and –2 °C during the first 90 min of the addition and allowed to rise to 0 °C during the last 30 min. The resulting solution was allowed to warm to room temperature with stirring over 1 h. Saturated aq. NaHCO₃ (2 dm³) was added to the reaction mixture followed by water (2 dm³) and the resulting biphasic system was stirred for 30 min; the layers were then separated. The organic phase was extracted with aq. NaHCO₃ (2.5 dm³) and then dried, (MgSO₄), filtered and evaporated under reduced pressure to give 11 (340 g, 98%) as a white solid, mp 129–134 °C (Found: C, 43.2; H, 4.2; N, 4.3. C₁₂H₁₄BrNO₃S requires C, 43.4; H, 4.25; N, 4.2%); $\nu_{\text{max}}(\text{KBr})/\text{cm}^{-1}$ 1735s (CO), 1340s and 1160s (SO₂); $\delta_{\text{H}}(\text{CDCl}_3)$ 2.45 (3 H, s, 4'-Me), 2.66 (1 H, dddd, J 14.8, 8.9, 5.7 and 1.2, 5-H_{ax}), 2.96 (1 H, ddd, J 14.8, 5.7 and 4.5, 5-H_{eq}), 3.25 (1 H, dddd, J 12.4, 8.9, 4.5 and 1.2, 6-H_{ax}), 3.36 (1 H, ddd, J 12.8, 8.4 and 1.2, 2-H_{ax}), 3.65 (1 H, dtd, J 12.4, 5.7 and 1.8, 6-H_{eq}), 3.98 (1 H, ddd, J 12.8, 5.1 and 1.8, 2-H_{eq}), 4.55 (1 H, ddd, J 8.4, 5.1 and 1.2, 3-H), 7.36 (2 H, d, J 8, 3'- and 5'-H) and 7.7 (2 H, d, J 8, 2'- and 6'-H); m/z 331 and 333 (M⁺).

3-Bromo-5-ethoxycarbonyl-1-(4'-methylphenylsulfonyl)azepan-4-one 12

To a solution of 11 (25.1 g, 75.6 mmol) in dichloromethane (822 cm³) at –5 °C under nitrogen, a solution of boron trifluoride–diethyl ether (9.96 cm³, 79.3 mmol) in dichloromethane (117 cm³) was added dropwise over 15 min, the temperature of the mixture being kept between –5 and –3 °C. The solution was stirred at –5 °C for 20 min after which a solution of ethyl diazoacetate (9.93 cm³, 94.4 mmol) in dichloromethane (117

cm³) was added dropwise to it during 20 min, the temperature of the solution being maintained between –5 and –2 °C during the addition. The solution was allowed to warm to room temperature for 90 min, after which it was diluted with water (234 cm³) and then stirred at room temperature for 25 min. The layers were separated and the organic layer was dried (Na₂SO₄), filtered and concentrated under reduced pressure to give a pale yellow oily solid which was crystallised from ethyl acetate to yield 12 (22.4 g, 71%) as a colourless solid, mp 165–167 °C (Found: C, 45.6; H, 4.75; N, 3.2. C₁₆H₂₀BrNO₅S requires C, 45.9; H, 4.8; N, 3.3%); $\nu_{\text{max}}(\text{KBr})/\text{cm}^{-1}$ 1740s and 1710s (CO), 1340s and 1160s (SO₂); $\delta_{\text{H}}(\text{CDCl}_3)$ 1.26 (3 H, t, CH₃CH₂O), 2.05 (1 H, dtd, J 14.7, 12.1 and 4.9, 6-H_b), 2.20 (1 H, dq, J 14.7 and 2.9, 6-H_a), 2.45 (3 H, s, 4'-Me), 2.81 (1 H, ddd, J 14.0, 12.1 and 2.9, 7-H_b), 3.07 (1 H, dd, J 15.3 and 11.0, 2-H_b), 4.00 (1 H, dddd, J 14.0, 4.9, 2.9 and 0.9, 7-H_b), 4.02 (1 H, dd, J 12.1 and 2.9, 5-H_a), 4.19 (2 H, q, CH₃CH₂O), 4.27 (1 H, m, J 15.3, 6.5 and 0.9, 2-H_a), 4.45 (1 H, dd, J 11.0 and 6.5, 3-H), 7.34 (2 H, d, J 8, 3'- and 5'-H) and 7.67 (2 H, d, J 8, 2'- and 6'-H); m/z 418 and 420 (MH⁺).

3-Bromo-1-(4'-methylphenylsulfonyl)azepan-4-one 13

To a suspension of 12 (45.4 g, 0.109 mol) in 1,4-dioxane (680 cm³) at 80 °C aq. HCl (3 mol dm^{–3}, 364 cm³) was added during 10 min. The resulting solution was heated at reflux for 7 h, cooled to room temperature and kept for 16 h. After this it was evaporated under reduced pressure to give a pale brown solid which was dissolved in ethyl acetate (650 cm³) and the solution washed with water (2 × 50 cm³) and brine (50 cm³) dried (Na₂SO₄), filtered, and evaporated under reduced pressure to afford a pale brown crystalline solid. This was recrystallised from diisopropyl ether to give 13 (33.9 g, 90%) as a colourless crystalline solid, mp 104–106 °C (Found: C, 45.2; H, 4.7; N, 4.0. C₁₃H₁₆BrNO₃S requires C, 45.1; H, 4.7; N, 4.05%); $\nu_{\text{max}}(\text{KBr})/\text{cm}^{-1}$ 1710s (CO), 1340s and 1160s (SO₂); $\delta_{\text{H}}(\text{CDCl}_3)$ 1.87 (1 H, m, 6-H_b), 1.96 (1 H, m, 6-H_a), 2.45 (3 H, s, 4'-Me), 2.56 (1 H, ddd, J 12.4, 6.8 and 2.6, 5-H_b), 2.77 (1 H, ddd, J 13.7, 11.7 and 3.4, 7-H_b), 2.88 (1 H, td, J 12.4 and 3.4, 5-H_a), 3.05 (1 H, dd, J 15.1 and 11.0, 2-H_b), 4.0 (1 H, dt, J 13.7 and 3.8, 7-H_a), 4.24 (1 H, ddd, J 15.1, 6.4 and 1.10, 2-H_a), 4.38 (1 H, dd, J 11.0 and 6.4, 3-H), 7.34 (2 H, d, J 8, 3'- and 5'-H) and 7.67 (2 H, d, J 8, 2'- and 6'-H); m/z 346 and 348 (MH⁺).

3-Azido-1-(4'-methylphenylsulfonyl)azepan-4-one 14

To a solution of 13 (33.9 g, 97.9 mmol) in dry DMF (750 cm³) under argon, acetic acid (11.2 cm³) and then sodium azide (12.7 g, 0.195 mol) were added. The resulting suspension was stirred at room temperature for 3.5 h and then kept at room temperature for 16 h. Upon dilution of the mixture with water (1.875 dm³) an oily solid separated and this was extracted with ethyl acetate (1 × 900 cm³, 3 × 300 cm³). The combined extracts were washed with brine (225 cm³) dried (Na₂SO₄), filtered and evaporated under reduced pressure to yield a yellow viscous liquid which crystallised when kept at room temperature. Recrystallisation of this from *tert*-butyl methyl ether gave 14 (21.9 g, 73%) as a white crystalline solid, mp 88 °C (Found: C, 50.3; H, 5.15; N, 17.8. C₁₃H₁₆N₄O₃S requires C, 50.6; H, 5.2; N, 18.1%); $\nu_{\text{max}}(\text{KBr})/\text{cm}^{-1}$ 2090s (N₃), 1710s (CO), 1340s and 1160s (SO₂); $\delta_{\text{H}}(\text{CDCl}_3)$ 1.89 (1 H, dddd, J 14.8, 8.2, 6.5 and 4.2, 6-H_b), 1.95 (1 H, m, 6-H_a), 2.45 (3 H, s, 4'-Me), 2.64 (1 H, ddd, J 13.8, 9 and 4.2, 5-H_b), 2.69 (1 H, ddd, J 13.8, 8.2 and 4.2, 5-H_a), 3.02 (1 H, ddd, J 13.7, 8.2 and 4.2, 7-H_b), 3.04 (1 H, dd, J 14.8 and 8.7, 2-H_a), 3.68 (1 H, dddd, J 13.7, 6.5, 4.2 and 1, 7-H_a), 3.78 (1 H, ddd, J 14.8, 5.3 and 1, 2-H_a), 4.21 (1 H, dd, J 8.7 and 5.3), 7.34 (2 H, d, J 8, 3'- and 5'-H) and 7.68 (2 H, d, J 8, 2'- and 6'-H).

trans-3-Azido-1-(4'-methylphenylsulfonyl)azepan-4-one 15 and cis-3-azido-1-(4'-methylphenylsulfonyl)azepan-4-one 16

To a suspension of 14 (21.8 g, 70.7 mmol) in ethanol (220 cm³)

at 0 °C sodium boranuide (2.67 g, 70.06 mmol) was added portionwise over 10 min. The resulting clear solution was stirred at 2–5 °C for 15 min after which it was cooled in an ice–water bath whilst water (670 cm³) was added to it followed by aq. HCl (1 mol dm^{–3}; 110 cm³), added dropwise over 10 min as the mixture was allowed to warm to ca. 20 °C. The mixture was then stirred at ca. 20 °C for 15 min after which it was further diluted with water (440 cm³) to give separation of a yellow oil which was extracted with ethyl acetate (1 × 440 cm³, 3 × 220 cm³). The combined extracts were washed with brine (2 × 55 cm³), dried (Na₂SO₄), filtered and evaporated under reduced pressure. The products were purified on SiO₂ eluting with *tert*-butyl–methyl ether–pentane (1:1, v/v to 7:3, v/v): to give 16 (13.5 g, 62%) as a white solid, mp 87–90 °C (Found: C, 50.15; H, 5.75; N, 18.15. C₁₃H₁₈N₄O₃S requires C, 50.31; H, 5.85; N, 18.05%); ν_{max} (KBr)/cm^{–1} 3510br(OH), 2940s(N₃), 1340s and 1160s(SO₂); δ_{H} (CDCl₃) 1.73 (2 H, m, 6-H), 1.98 (2 H, m, 5-H), 2.05 (1 H, dd, J 3.6 and 1.2, -OH), 2.45 (3 H, s, 4'-Me), 2.93 (1 H, dt, J 12.4 and 5.9, 7-H_b), 3.08 (1 H, dd, J 14.4 and 9.6, 2-H_b), 3.61 (2 H, m, 2-H_a and 7-H_a), 3.78 (1 H, ddd, J 9.6, 3.9 and 3.3, 3-H), 4.1 (1 H, br m, 4-H), 7.34 (2 H, d, J 8, 3'- and 5'-H) and 7.68 (2 H, d, J 8, 2'- and 6'-H); and 15 (5.7 g, 26%) as a white solid, mp 67–70 °C (Found: C, 50.2; H, 5.8; N, 18.1. C₁₃H₁₈N₄O₃S requires C, 50.3; H, 5.85; N, 18.05%); ν_{max} (KBr)/cm^{–1} 3510br(OH), 2940s(N₃), 1340s and 1160s(SO₂); δ_{H} (CDCl₃) 1.54 (1 H, m, 6-H_b), 1.78 (1 H, m, 6-H_a), 1.96 (2 H, m, 5-H), 2.22 (1 H, d, J 3.9, OH), 2.45 (3 H, s, 4'-Me), 2.82 (1 H, dd, J 14.8 and 9, 2-H_b), 3.01 (1 H, ddd, J 12.2, 6.5 and 3.5, 7-H_b), 3.48 (2 H, m, 4-H and 7-H_a), 3.53 (1 H, dd, J 9 and 3.6, 3-H), 3.66 (1 H, ddd, J 14.8, 3.6 and 0.9, 2-H_a), 7.34 (2 H, d, J 8, 3'- and 5'-H) and 7.68 (2 H, d, J 8, 2'- and 6'-H).

trans-3-Azido-1-(4'-methylphenylsulfonyl)-4-(4"-nitrobenzoyloxy)azepane 17

To a solution of triphenylphosphine (17.6 g, 67.1 mmol) in dry THF (330 cm³) at 5 °C, DIAD (diisopropylazodicarboxylate; 13.9 cm³, 67.1 mmol) was added dropwise during 10 min while the temperature of the mixture was kept at 5–10 °C. A thick cream suspension formed which was stirred for a further 15 min and to this 4-nitrobenzoic acid (11.2 g, 67 mmol) and 16 (13.4 g, 43.2 mmol) were added. The resulting yellow solution was allowed to warm to room temperature as it was stirred for 1 h and then kept at room temperature for 16 h. After this the mixture was evaporated under reduced pressure and the residue stirred with *tert*-butyl methyl ether (170 cm³) to yield a cream solid. The product was crystallised from *tert*-butyl methyl ether to give 17 (16.9 g, 85%) as colourless crystals, mp 164–168 °C (Found: C, 52.3; H, 4.6; N, 15.1. C₂₀H₂₁N₃O₆S requires C, 52.3; H, 4.6; N, 15.2%); ν_{max} (KBr)/cm^{–1} 2105s(N₃), 1720s(CO), 1530s and 1350w(NO₂), 1330s and 1160s(SO₂); δ_{H} (CDCl₃) 1.70 (1 H, m, 6-H_b), 2.06 (3 H, m, 5-H and 6-H_a), 2.45 (3 H, s, 4'-Me), 2.82 (1 H, dd, J 15 and 10, 2-H_b), 2.97 (1 H, ddd, J 12.3, 6.6 and 3.5, 7-H_b), 3.64 (1 H, ddd, J 12.3, 9.8 and 6.6, 7-H_a), 3.76 (1 H, ddd, J 15, 4.2 and 1, 2-H_a), 4.02 (1 H, ddd, J 10, 8.4 and 4.2, 3-H), 5.0 (1 H, ddd, J 10, 8.4 and 2.9, 4-H), 7.34 (2 H, d, J 8, 3'- and 5'-H), 7.7 (2 H, d, J 8, 2'- and 6'-H), 8.25 (2 H, d, J 8, 3'- and 5'-H) and 8.33 (2 H, d, J 8, 2'- and 6'-H).

Conversion of *trans*-3-azido-1-(4'-methylphenylsulfonyl)-4-(4"-nitrobenzoyloxy)azepane 17 into *trans*-3-azido-1-(4'-methylphenylsulfonyl)azepan-4-ol 15

To a suspension of 17 (16.9 g, 36.8 mmol) in methanol (750 cm³) and 1,4-dioxane (190 cm³) was added aq. NaOH (2% w/v; 150 cm³) at room temperature. The suspension was stirred at room temperature for 5 h; a clear solution was obtained after 3 h. The solution was kept at room temperature for 16 h, after which it was evaporated under reduced pressure to give an oily residue which was stirred with ethyl acetate (460 cm³) and water (460

cm³). The layers were separated and the aqueous layer was extracted with ethyl acetate (3 × 230 cm³). The combined organic layer and extracts were washed with brine (2 × 50 cm³), dried (Na₂SO₄), filtered and concentrated under reduced pressure to give a viscous liquid, which crystallised with time to give 15 (11.3 g, 99%) mp 66–70 °C, identical with 15 as prepared above.

trans-3-Amino-1-(4'-methylphenylsulfonyl)azepan-4-ol 18

To a solution of 15 (16.9 g, 54.5 mmol) in dry THF (270 cm³) at 0 °C under nitrogen, a solution of lithium aluminium hydride in THF (1 mol dm^{–3}; 27.3 cm³, 27.3 mmol) was added dropwise during 10 min, while the temperature of the mixture was kept at 0–2 °C. After the mixture had been allowed to warm to room temperature it was stirred for 45 min. Water (30 cm³) was then added dropwise during 10 min to the mixture with cooling (ice–water bath) followed by aq. Na₂CO₃ (275 cm³; 20% w/v). The mixture was diluted with water (2 dm³) and extracted with ethyl acetate (1 × 1 dm³, 3 × 450 cm³) and the combined extracts were washed with brine (2 × 150 cm³), dried (Na₂SO₄), filtered and evaporated under reduced pressure to give a colourless oil which crystallised with time. The product was recrystallised from *tert*-butyl methyl ether to give 18 (13.1 g, 85%) as colourless crystals, mp 93–96 °C (Found: C, 54.8; H, 7.05; N, 9.6. C₁₃H₂₀N₂O₃S requires C, 54.9; H, 7.1; N, 9.85%); ν_{max} (KBr)/cm^{–1} 3360br(OH), 3100br and 2930br(NH₂), 1330s and 1150s(SO₂); δ_{H} (CDCl₃) 1.64 (2 H, m, 6-H), 1.94 (2 H, m, 5-H), 2.45 (3 H, s, 4'-Me), 2.76 (1 H, td, J 8.1, 8.1 and 3.5, 3-H), 2.86 (1 H, dd, J 14.2 and 8.1, 2-H_b), 3.18 (1 H, ddd, J 12.6, 6.1 and 4.4, 7-H_b), 3.22 (1 H, m, 7-H_a), 3.29 (1 H, ddd, J 9.8, 8.1 and 2.4, 4-H), 3.49 (1 H, ddd, J 14.2, 3.5 and 0.6, 2-H_a), 7.31 (2 H, d, J 8, 3'- and 5'-H) and 7.66 (2 H, d, J 8, 2'- and 6'-H); *m/z* 285 (MH⁺).

trans-3-Aminoazepan-4-ol dihydrobromide 19

A suspension of 18 (13.0 g, 45.7 mmol) in aq. HBr (48%; 130 cm³) was stirred and heated to reflux at which temperature the resulting yellow solution was kept for 3 h. The solution was then cooled to room temperature and kept for 16 h after which it was diluted with ice–water (260 cm³) and then extracted with *tert*-butyl methyl ether (1 × 160 cm³, 2 × 40 cm³). The aqueous layer was concentrated under reduced pressure to give an oily yellow solid which was crystallised from ethanol to yield 19 (9.09 g, 68%) as a white crystalline solid, mp 215 °C (decomp.) (Found: C, 24.5; H, 5.6; N, 9.4. C₆H₁₆Br₂N₂O requires C, 24.7; H, 5.5; N, 9.4%); ν_{max} (KBr)/cm^{–1} 3420br, 3230br and 2900br; δ_{H} (CD₃SOCD₃) 1.6–1.65 (3 H, m), 1.79–1.81 (1 H, m), 1.95–2.05 (1 H, m), 3.05–3.2 (3 H, m), 3.65 (1 H, br), 5.75 (1 H, br), 8.18 (3 H, br) and 9.1 (1 H, br).

trans-3-Amino-1-benzoyloxycarbonylazepan-4-ol 20

To a stirred suspension of 19 (9.01 g, 30.9 mmol) in dry dichloromethane (230 cm³) at room temperature under nitrogen, triethylamine (21.5 cm³) was added followed by 18-crown-6 (16.3 g, 61.7 mmol). The mixture was stirred at room temperature for 20 min and the resulting clear solution cooled to 5 °C and benzyl chloroformate (4.97 cm³, 33.9 mmol) was added dropwise during 10 min. The mixture was stirred at room temperature for 5 h, allowed to stand at room temperature for 16 h and then concentrated under reduced pressure to give a pale yellow oily solid which was stirred with ethyl acetate (330 cm³) and aq. Na₂CO₃ (10% w/v; 220 cm³). The layers were separated and the aqueous layer was extracted with ethyl acetate (3 × 110 cm³). The combined organic extracts were washed with aq. Na₂CO₃ (10% w/v; 110 cm³) and saturated aq. KBr (2 × 55 cm³), dried (Na₂SO₄), filtered and the mixture was concentrated under reduced pressure to give a pale yellow oily solid. Crystallisation from *tert*-butyl methyl ether gave 20

(7.19 g, 88%) as a colourless crystalline solid, mp 78–80 °C (Found: C, 63.7; H, 7.8; N, 10.7. $C_{14}H_{20}N_2O_3$ requires C, 63.6; H, 7.6; N, 10.6%); ν_{max} (KBr)/cm⁻¹ 3350br, 3270br and 2900br, 1700s(CO); δ_H (CDCl₃) 1.48 (1 H, m, 6-H_b), 1.61 (1 H, m, 6-H_a), 1.96 (2 H, m, 5-H), 2.66 (4 H, dt, J 9.9 and 3.7, 3-H), 2.70 (4 H, dt, J 9.9 and 3.7, 3-H), 2.87 (4 H, dd, J 14.4 and 9, 2-H_b), 2.97 (4 H, dd, J 14.4 and 9, 2-H_b), 3.22 (1 1/2 H, m, 7-H_b and 4-H), 3.33 (2 H, ddd, J 13.6, 5.9 and 4.3, 7-H_b), 3.53 (4 H, ddd, J 13.6, 9.9 and 5.9, 7-H_a), 3.66 (4 H, ddd, J 13.6, 9.9 and 5.9, 7-H_a), 3.71 (1 H, dd, J 14.5 and 3.8, 2-H_a), 3.79 (4 H, dd, J 14.5 and 3.8, 2-H_a), 5.12, 5.17 (AB system, 2 H, J 12, PhCH₂) and 7.28–7.4 (5 H, m, Ar); m/z 265 (MH⁺).

trans-3-[4-(Benzylxy)benzamido]-1-benzylxycarbonylazepan-4-ol 21

To a stirred suspension of 20 (3.09 g, 11 mmol) in dichloromethane (60 cm³) at 0 °C, triethylamine (2.28 cm³, 16.4 mmol) was added. While the temperature of the mixture was maintained at ca. 0 °C, 4-(benzylxy)benzoyl chloride (3.02 g, 11 mmol) in dichloromethane (30 cm³) was added dropwise after which the mixture was stirred at ca. 0 °C for a further 3 h. The mixture was then evaporated under reduced pressure and the residue dissolved in ethyl acetate (500 cm³). The resulting solution was washed with water (2 × 60 cm³) and brine (1 × 60 cm³), dried (Na₂SO₄), filtered and evaporated under reduced pressure to yield a viscous yellow oil. Chromatography of the oil on silica, eluting with *tert*-butyl methyl ether gave a colourless viscous oil, crystallisation of which from diisopropyl ether gave 21 (3.29 g, 63%) as a colourless solid, mp 132–134 °C (Found: C, 70.8; H, 6.4; N, 5.8. $C_{28}H_{30}N_2O_3$ requires C, 70.9; H, 6.4; N, 5.9%); ν_{max} (KBr)/cm⁻¹ 3330br (OH), 2950br (NH) and 1700s(CO); δ_H (CDCl₃) 1.66 (1 H, m, 6-H_b), 1.78–2.0 (3 H, m, 5- and 6-H_a), 2.78 (1 H, ddd, J 14, 13 and 4, 7-H_b), 3.35 (1 H, dd, J 15 and 5, 2-H_b), 3.78 (1 H, ddd, J 10, 6 and 2, 4-H), 4.07–4.23 (3 H, m, 2-H_a, 3-H and 7-H_a), 5.12 (2 H, s, OCH₂Ph), 5.14, 5.21 (AB system, 2 H, J 12, PhCH₂), 5.46 (1 H, br, s, OH), 7.02 (2 H, d, J 8, Ar), 7.28–7.47 (10 H, m, Ar), 7.82 (2 H, d, J 8, Ar) and 8.83 (1 H, br d, NH); m/z 475 (MH⁺). Spectroscopic data of 21 were identical with those reported.⁸

Separation of the enantiomers of 21

To a solution of 21 (170 mg, 0.36 mmol), triethylamine (0.1 cm³, 0.72 mmol) and DMAP (4-dimethylaminopyridine; 44 mg, 0.36 mmol) in dichloromethane (5 cm³), the acid chloride derived from (S)-(–)-MTPA [α -methoxy- α -(trifluoromethyl)phenylacetic acid; 113 mg, 0.45 mmol] in dichloromethane (2 cm³) was added dropwise at room temperature. The mixture was stirred at room temperature for 16 h after which it was evaporated under reduced pressure. The residue was chromatographed on silica eluting with ethyl acetate–pentane (1:3, v/v) to give the enantiomerically pure Mosher's esters 22a (120 mg, 48%); m/z 691 (MH⁺), and 22b (120 mg, 48%); m/z 691 (MH⁺). Both 22a and 22b appear as a ca. 3:1 rotameric mixture by ¹H NMR. Separate solutions of both 22a (120 mg, 0.17 mmol) and 22b (120 mg, 0.17 mmol) in methanol (5 cm³) were stirred and treated with aq. KOH (1 mol dm⁻³; 2 cm³). Each reaction mixture was stirred at room temperature for 16 h and then diluted with diethyl ether (30 cm³) and water (20 cm³). The organic layer was separated and evaporated under reduced pressure to yield a colourless gum, treatment of which with diisopropyl ether gave, as white solids, 21a (80 mg, 100%), mp 113–114 °C, and 21b (80 mg, 100%), mp 116–118 °C. Both 21a and 21b were identical spectroscopically with 21. The enantiomeric purity of 21a and 21b was assessed relative to racemic 21 using chiral chromatography on a Chiral Pack AD column (4.6 mm × 250 mm) eluting with ethanol–heptane (15:85, v/v) which showed 21a and 21b both to be >99% enantiomerically pure.

Protected balanol 23

To a stirred solution of 10 (180 mg, 0.265 mmol), triethylamine (0.74 cm³, 5.3 mmol) and DMAP (16.4 mg, 0.133 mmol) in dichloromethane (9 cm³), 2-chloro-1-methylpyridinium iodide (88 mg, 0.345 mmol) was added and the mixture stirred at room temperature for 1 h. Compound 21 (125 mg, 0.26 mmol) was added to the mixture which was then, stirred at room temperature for 20 h and finally heated at reflux for 2 h. After being cooled to room temperature the mixture was evaporated under reduced pressure and the residue dissolved in ethyl acetate (20 cm³). The resulting solution was washed with aq. NaHCO₃ (5%; 2 × 50 cm³), dried (MgSO₄), filtered and evaporated under reduced pressure. Purification of the residue on silica eluting with ethyl acetate–dichloromethane (5:95, v/v) gave 23 (0.11 g, 37%) as a glass; m/z 1135 (MH⁺); δ_H (CDCl₃) as a ca. 3:1 rotameric mixture 1.6–2.1 (4 H, m), 2.9 (1 H, m), 3.4 (1 H, m), 4.08–4.16 (2 H, m), 4.67 (2 H, s, OCH₂Ph), 4.7–4.88 (1 H, m), 4.82, 4.85 (2 H, AB system, J 12, OCH₂Ph), 5.5–5.15 (1 H, m), 5.04 (2 H, s, OCH₂Ph), 5.1 (4 H, s, OCH₂Ph), 5.26 (2 H, s, OCH₂Ph), 6.82 (2 H, d, J 8, Ar), 6.88–6.95 (4 H, m, Ar), 7.0–7.45 (32 H, m, Ar), 7.72 and 7.79 (ca. 3:1, 2 H, d, J 8, Ar). Spectroscopic data were identical with those reported;⁸ the presence of rotamers precludes the exact assignment of all resonances.

Banol 1

A solution of 23 (106 mg, 0.093 mmol) in ethyl acetate–acetic acid–water (19 cm³, 16:2:1, v/v) was treated with palladium black (10.6 mg) under an atmosphere of hydrogen. The reaction mixture was initially warmed to 50 °C and then stirred at room temperature for 4 h. After this the mixture was filtered and fresh catalyst (10.6 mg) added, to the filtrate; the reaction mixture was then placed under a fresh atmosphere of hydrogen and stirred at room temperature for 24 h. The catalyst was filtered off and washed with a solvent mixture (16 cm³) as used for the reaction. The mixture was concentrated and the residue was chromatographed on a Dynamax C₁₈ column (20 × 300 mm) eluting with acetonitrile–water–trifluoroacetic acid (20:80:0.1, v/v) to give 1 (27 mg, 53%) as a light yellow powder; m/z 551 (MH⁺); δ_H (CD₃OD) 1.84–2.12 (4 H, br m, 5- and 6-H), 3.42–2.98 (4 H, br m, 2- and 7-H), 4.32 (1 H, br m, 3-H), 5.29 (1 H, m, 4-H), 6.76 (2 H, d, J 8.7, 4'- and 6'-H), 6.80 (1 H, d, J 7.8, 11'-H), 6.92 (2 H, s, 3'- and 7'-H), 7.17 (1 H, t, J 7.8, 12'-H), 7.25 (1 H, d, J 7.8, 13'-H) and 7.60 (2 H, d, J 8.7, 3', 7'-H). Spectroscopic data were identical with those reported.⁸

X-Ray crystal structure determination of *trans*-3-azido-1-(4'-methylphenylsulfonyl)hexahydroazepan-4-ol 15

A single crystal of compound 15 (from *tert*-butyl methyl ether, approximate size 0.18 × 0.28 × 0.16 mm), mounted in a Lindemann tube, was used for X-ray data collection.

Crystal data. $C_{13}H_{18}N_4O_3S$, $M = 310.37$, colourless prisms, orthorhombic, space group $Pn2_1a$, $a = 8.7030(9)$, $b = 11.3280(13)$, $c = 15.097(2)$ Å (by least squares refinement of the setting angles for 250 reflections within $\theta = 2.25$ –27.16 °), $V = 1488.4(3)$ Å³, $Z = 4$, $D_c = 1.385$ g cm⁻³, $T = 120$ K, $\mu(\text{MoK}\alpha) = 2.33$ cm⁻¹, $F(000) = 656$.

Data collection, structure solution and refinement. Data were collected on a FAST TV Area detector diffractometer following previously described methods.¹⁵ From the ranges scanned, 6059 data were recorded (2.25 < θ < 27.16 °; index ranges $-10 \leq h \leq 10$, $-13 \leq k \leq 9$, $-18 \leq l \leq 10$) and merged to give 2618 unique [R (int) = 0.0532]. The structure was solved by direct methods¹⁶ and refined on F_0^2 by full matrix least squares¹⁷ using all unique data corrected for Lorentz and polarisation factors. All non-hydrogen atoms were anisotropic. The hydrogen atoms were inserted in idealised positions with U_{iso} set at 1.5 times U_{eq} of the parent. The weighting scheme

used was $w = 1/[\sigma^2(F_o)^2 + (0.0612P)^2]$, where $P = [\max(F_o)^2 + 2(F_o)^2]/3$; this gave satisfactory agreement analysis. Final R_1 (on F) and R_{w2} (on F_o^2) values were 0.0473 and 0.1067 for all 2616 data and 191 parameters. The corresponding R values were 0.0424 and 0.1008 for 2323 data with $I > 2\sigma(I)$. Sources of scattering factors are given in ref. 17.

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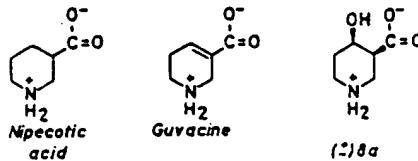
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Inhibitors of GABA Uptake. Syntheses and ^1H NMR Spectroscopic Investigations of Guvacine, (3RS,4SR)-4-Hydroxypiperidine-3-carboxylic Acid, and Related Compounds

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The syntheses of (3RS,4SR)-4-hydroxypiperidine-3-carboxylic acid (*8a*) and guvacine (1,2,5,6-tetrahydropyridine-3-carboxylic acid hydrobromide (*9a*), both potent inhibitors of γ -aminobutyric acid (GABA) uptake, are described. Furthermore (3RS,4SR,5SR)- and (3RS,4SR,5RS)-4-hydroxy-5-methyl(piperidine-3-carboxylic acids (*8c*) and (*8d*) and the guvacine analogues (RS)-5-methyl-1,2,5,6-tetrahydropyridine-3-carboxylic acid (*10*) and 2,5,6,7-tetrahydro-1*H*-azepine-3-carboxylic acid and (1*R*,5*S*)-(-)-2-nortropene-2-carboxylic acid hydrobromides (*14*) and (*18*) have been synthesized. The compounds *8a,c,d*, *9a*, *10*, and *14* were prepared via catalytic hydrogenation of cyclic β -oxoesters and appropriate acid treatments of the intermediate β -hydroxy esters. Demethylation of ecgonine (*16*) followed by acid catalyzed hydrolysis and elimination reactions gave *18*. (RS)-Perhydroazepine-3-carboxylic acid and (1*R*,2*R*,5*R*)-(+)-nortropane-2-carboxylic acid hydrobromides (*15*) and (*19*) were obtained by catalytic hydrogenation of *14* and *18*, respectively. The relative stereochemistry of *8a,c,d*, *12*, *13*, and *19* was established by 270 MHz ^1H NMR spectroscopy. The relationship between structure and potency as inhibitors of GABA uptake of *8a,c,d*, *9a*, *10*, *14*, *15*, *18*, and *19* is discussed.



Scheme 1.

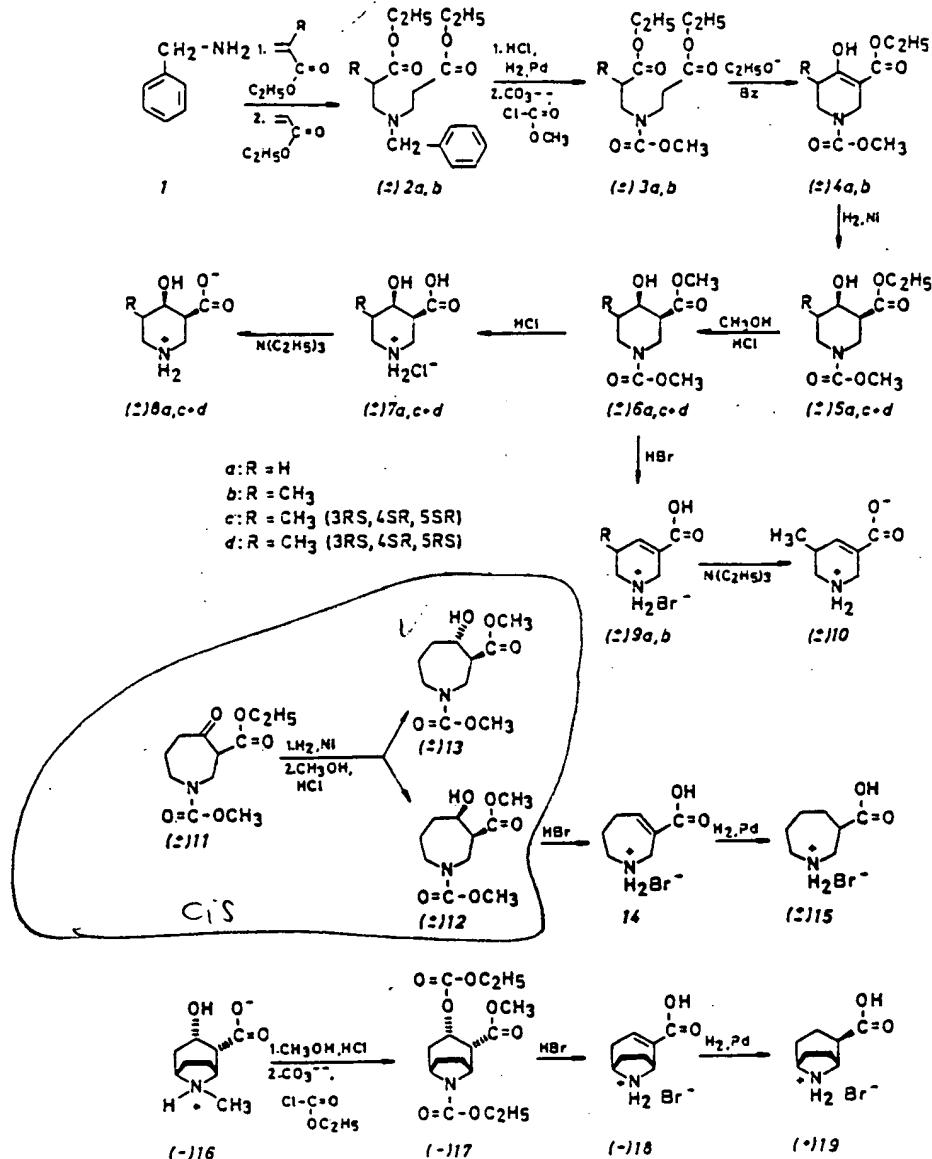
Nipecotic acid (piperidine-3-carboxylic acid),¹⁻³ guvacine (1,2,5,6-tetrahydropyridine-3-carboxy-

lic acid),⁴ and (3RS,4SR)-4-hydroxypiperidine-3-carboxylic acid (*8a*)⁵ (Scheme 1) are potent substrate-competitive inhibitors of the neuronal γ -aminobutyric acid (GABA) uptake process. The concentrations of (*R*)-(-)-nipecotic acid, guvacine, and compound *8a* required for 50 % inhibition of GABA uptake (IC_{50} values) are 5 μM ,⁵ 8 μM ,⁴ and 12 μM ,⁵ respectively. These compounds seem to combine with the GABA transport carrier and penetrate the tissue.²⁻⁴ Such compounds have pharmacological interest and may be useful tools for the study of the GABA transport carrier. However, molecular manipulations of these amino acids result in compounds with considerably reduced potency as inhibitors of GABA uptake, the IC_{50} values of *8c*, *8d*, *10*, and *15* being 260,⁶ 349,⁶ 547,⁸ and 502 μM ,⁶ respectively. Compound *14*⁶ and the conformationally rigid amino acids *18*⁶ and *19*⁶ are almost inactive. These findings demonstrate a pronounced substrate specificity of the GABA transport carrier.

This paper describes the syntheses of *8a*, its 5-methyl derivatives *8c* and *8d*, and the guvacine and nipecotic acid analogues *10*, *14*, *18*, *15*, and *19*. A convenient method for the preparation of guvacine hydrobromide (*9a*) has been developed. Guvacine hydrochloride has previously been synthesized on a very small scale.^{1,6}

The β -oxoester *4b* was the only detectable product after Dieckmann cyclization of the unsymmetrical diester *3b* (Scheme 2). High pressure hydrogenation of *4a,b* gave the 3,4-cis-4-hydroxynipecotic acid derivatives *5a* and

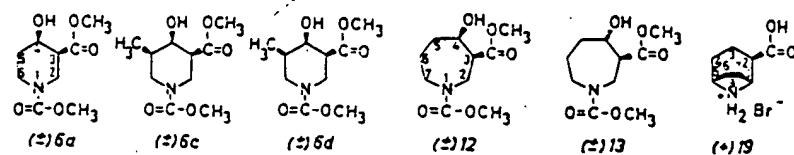
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Scheme 2.

a separable mixture of *5c* and *5d*, respectively. The ethyl esters *5a,c,d* were transformed into the corresponding methyl esters in order to facilitate the analysis of the 270 MHz ^1H NMR spectra. Reduction of the β -oxoester *11*, however, gave a separable mixture of the *cis*- and *trans*- β -hydroxy esters *12* and *13* (Scheme 2).

Appropriate treatments of *6a,c,d* with hydrochloric acid gave the hydroxy amino acid chlorides *7a,c,d*, whereas the compounds *9a,b* were synthesized by prolonged treatments of *6a* and *6c* with hydrobromic acid. *N*-Demethylation of the methyl ester of ecgonine [(*1R,2R,3S,5S*)-(-)-3-hydroxytropane-2-carboxylic acid].

Table 1. Some chemical shifts (δ) and coupling constants (Hz) from the 270 MHz ^1H NMR spectra of *6a,c,d*, *12*, *13*, and *19*.^a C_2H_5
=OH₃ C_2H_5
=O

	(\pm) <i>6a</i>	(\pm) <i>6c</i>	(\pm) <i>6d</i>	(\pm) <i>12</i>	(\pm) <i>13</i>	($+$) <i>19</i>
δ_{1e}	—	—	—	—	—	4.15
δ_{2a}	3.43	3.19	ca. 3.7	3.38	3.17, 3.22	2.88
δ_{2c}	4.05	4.28	ca. 3.7	3.92, 4.07	3.79, 3.96	
δ_{3a}	2.64	2.58	2.81	2.71, 2.78	2.60, 2.66	
δ_{4a}	—	—	—	—	3.91	
δ_{4e}	4.32	4.17	3.79	4.32	—	
δ_{5a}	1.68	1.68	—	1.53	1.62	
δ_{5e}	1.84	—	1.98	2.06	1.90	3.93
δ_{6a}	3.30	2.82	3.23	1.69	1.62	
δ_{6e}	3.79	3.80	3.59	2.06	2.00	
δ_{7a}	—	—	—	3.20, 3.25	3.32	
δ_{7e}	—	—	—	3.64	3.56	
$\delta_{\text{C}-\text{COOCH}_3}$	3.68	3.69	3.69	3.71, 3.72	3.69	
$\delta_{\text{N}-\text{COOCH}_3}$	3.71	3.73	3.72	3.74, 3.75	3.73	
$\delta_{\text{C}-\text{CH}_3}$	—	0.99	0.98	—	—	
J_{1e2a}	—	—	—	—	—	2.2
J_{1e2c}	—13.2	—13.0	—	—14.5	—14.8	
J_{2a2a}	10.7	12.0	10.0 ^b	10.2	9.0	
J_{2a2c}	4.5	2.0	2.8 ^b	3.0	3.75	
J_{3a2e}	2.5	4.4	3.4	3.0	—	
J_{3a2a}	—	—	—	—	9.0	
J_{3a2c}	—	—	—	—	2	
J_{4a2a}	—	—	—	—	9	
J_{4a2c}	—	—	—	—	—	
J_{5a2a}	3.0	ca. 4	—	—	—	
J_{5a2c}	4.4	—	5.3	—	—	
J_{6a2e}	—13.6	—	—	—	—	
J_{6a2a}	11.7	12.0	—	—	—	
J_{6a2c}	4.8	ca. 2	—	—	—	
J_{7a2a}	3.2	—	5.4	—	—	
J_{7a2c}	3.4	—	3.4	—	—	
J_{7a2e}	—13.4	—13.0	—13.6	—	—	
J_{7a2e}	—	—	—	—14	—	

^a The spectra of *6a,c,d*, *12*, and *13* were recorded in CDCl_3 and that of *19* in D_2O solutions. ^b Only the sum of J_{2a2a} and J_{2c2a} is precisely determined.

(16) by a modified von Braun procedure⁹ followed by acid hydrolysis and dehydration of the intermediate *17* (Scheme 2) gave the rigid guvacine analogue *18*. Hydrogenation of *18* proceeded stereospecifically, *19* being the only detectable product.

The structure elucidation of the new products *2b-4b*, *5-10*, *12-15*, and *17-19* were based on elemental analyses, IR and ^1H NMR spec-

troscopy, in the cases of *9a,b*, *10*, *14*, and *18* supported by UV spectroscopy. In tetrachloromethane solution the β -oxoester *4b* (Scheme 2) exists in the enol-form as established by ^1H NMR spectroscopy. The relative configurations of *5-8*, *12*, *13*, and *19* were established by analysis of the 270 MHz ^1H NMR spectra of the methyl esters *6a,c,d*, *12*, and *13* and compound *19*, respectively.

with hydro-
amino acid
compounds *9a,b*
treatments of
N-Demethyl-
ine [(1*R*,2*R*,
3*S*)-*3*-*oxo* acid]

(1978) No. 5

Acta Chem. Scand. B 32 (1978) No. 5

In simple piperidine derivatives the equatorial proton on C(2) is found downfield from its axial counterpart.¹⁰ The coupling constants between the C(3) proton and the two C(2) protons are typical for equatorial-axial and axial-axial configurations of these protons in *6a*. This is consistent with a predominantly equatorial orientation of the C(3) methoxycarbonyl group (Table 1). Furthermore the coupling constant for the C(3) and C(4) protons unequivocally indicates axial-equatorial orientation of the protons concerned and therefore a 3,4-cis configuration of *6a*. An analysis of the mutual coupling constants for the C(4) and C(5) protons supports this assignment. The vicinal coupling constants used in this assignment are in general agreement with those found in most 6-membered rings.¹¹ The geminal coupling constants found parallel data previously found in other piperidines.¹² The coupling patterns of the C(2), C(3), and C(4) protons in *6c* are very similar to those of the corresponding system in *6a* establishing equatorial and axial positions of the substituents at C(3) and C(4), respectively. The coupling constants for the C(5) proton indicate an equatorial orientation of the C(5) methyl group. These findings together are in agreement with the depicted relative configuration of *6c*. In *6d* the mutual coupling constants for the C(6) and C(5) protons indicate an axial orientation of the C(5) methyl group. A further analysis of the C(5), C(4), and C(3) proton coupling patterns reveals axial and equatorial orientations of the hydroxy and C(3) methoxycarbonyl groups, respectively.

In previous studies of piperidines¹⁰ it has been suggested that the chemical shift difference between axial and equatorial protons on C(2) and C(6) ($\Delta\delta$) would indicate the extent to which the nitrogen lone pair influences the two protons. The differences $\Delta\delta(H_{2a},H_{2e})$ and $\Delta\delta(H_{5a},H_{5e})$ found in *6a* would be consistent with the dominating contribution from the equatorial-axial orientation of the C(3) and C(4) substituents of *6a*. The $\Delta\delta$ values found for *6c* are larger, indicating an increased fixation in an equatorial-axial-equatorial arrangement of the C(3), C(4), and C(5) substituents. This tendency is reflected in the values of the axial-axial coupling constants, where $J_{2aa} = 12.0$ Hz would be consistent with an almost locked conformation. In *6d* the spectrum is quite

different. The two protons at C(2) are almost coinciding in chemical shift at a value of ca. 3.7 ppm, the lack of accuracy arising from the near coincidence of the signal with those of the methoxy groups. The chemical shift difference between the two protons at C(2) can be judged from the triplet structure of the C(3) proton in the ABX pattern formed by the three protons at C(2) and C(3). The vanishingly small chemical shift difference on C(2) and the reduced chemical shift difference on C(6) as well as the smaller value of J_{2aa} in *6d* (10.0 Hz) all point to a conformational equilibrium where equatorial-axial-axial orientations of the C(3), C(4), and C(5) substituents represent the preferred but not the exclusive conformation.

The 270 MHz ¹H NMR spectra of *12* and *13* were analysed as far as the complexity of the spectra permitted. Interpretation of the spectra, recorded at 293 K, was complicated by the fact that both molecules gave rise to two sets of signals due to hindered rotation of the urethane group. This is reflected in dual values for several chemical shifts while the coupling constants determined in both conformers were identical. The coupling constants clearly indicate that the arrangement of the substituents in position 3 and 4 are equatorial-axial in *12* and equatorial-equatorial in *13*. The observed values for chemical shifts and coupling constants are in general agreement with the data given for *6a*. In the spectrum of *19* a merging and very complex pattern of signals from the C(3), C(4), C(6), and C(7) protons is observed. It is not possible to establish the orientation of the carboxyl group of *19* on the basis of the coupling constant for the equatorial C(1) proton and the proton on C(2). However, the total width of the pattern originating in the C(2) proton reveals the existence of an axial-axial coupling between this proton and the axially oriented C(3) proton thus indicating an equatorial orientation of the carboxyl group.

EXPERIMENTAL

Unless otherwise stated, the determination of melting points and elemental analyses and the recording of IR, UV, and 60 MHz ¹H NMR spectra were accomplished as described in a previous paper.¹³ ¹H NMR spectra of compounds dissolved in D₂O were recorded by using sodium 3-(trimethylsilyl)propanesulfonate as an in-

(2) are almost a value of ca. rising from the those of the shift difference can be judged C(3) proton in three protons small chemical reduced chemi- as the smaller point to a con- quatorial-axial- C(4), and C(5) erred but not

of 12 and 13 complexity of the of the spectra, ed by the fact to two sets of of the urethane tues for several ling constants were identical. indicate that nts in position and equatorial- ed values for nstants are in given for 6a. ring and very om the C(3), observed. It is ntation of the of the coupling proton and the total width of C(2) proton axial coupling nally oriented uatorial orien-

ternal standard. The 270 MHz ^1H NMR spectra were obtained on a Bruker HX 270 S instrument operating at 293 and 303 K. Fourier transform method was used to obtain the spectra with a spectral width of 1500 Hz using 32 K data points. The detection was quadrature detection. Homodecoupling was used to verify the interpretations of the spectra and provide starting parameters for the analyses of the spectra. Nitrogen decoupling was possible using a Bruker probehead equipped with an additional nitrogen decoupling coil. The frequency was derived from a Bruker synthesizer model BS 100 and fed to the amplifier of the decoupling channel through a band pass filter before entering the probe. The decoupling power was 5 watt. The decoupling frequency used was 19 506 582 Hz. The proton spectra were analyzed using the program SIMEQ and a Varian 620/i computer. Due to the large chemical shift differences at 270 MHz, it is permissible to subdivide the total spin system into several subsystems containing typically 3 to 6 nuclei. After the initial assignment the spectra were iterated to a best fit using the ITRCAL program on a Nicolet 1130 computer with 80 K memory. Optical rotations were measured on a Perkin-Elmer polarimeter model 141. Thin-layer chromatography (TLC) and column chromatography (CC) were accomplished by using silica gel F_{254} plates (Merck) and silica gel (Woelm 0.063–0.1 mm), respectively. Columns were developed by stepwise gradient elution. The pK_A values were determined as described in a previous paper.¹⁴

(RS)-Ethyl N-benzyl-N-(2-ethoxycarbonylethyl)-2-methyl-3-aminopropionate (2b). To a solution of 1 (107 g; 1 mol) in ethanol (200 ml) was added ethyl methacrylate (137 g; 1.2 mol). The mixture was refluxed for 24 h. The solvent removed in *vacuo*, and the residue distilled to give (RS)-ethyl N-benzyl-2-methyl-3-aminopropionate (149.9 g; 68 %), b.p. 131–133 °C/80 Pa. Anal. $\text{C}_{13}\text{H}_{17}\text{NO}_3$; C, H, N. IR (film): 3330 (w), 3030 (w), 2970–2820 (m), 1725 (s), 1495 (m), 1450 (m) cm^{-1} . ^1H NMR (60 MHz, CCl_4): δ 7.17 (5 H, s), 3.98 (2 H, q, J 7 Hz), 3.63 (2 H, s), 3.0–2.1 (3 H, m), 1.42 (1 H, s), 1.33–0.97 (6 H, m).

A solution of (RS)-ethyl N-benzyl-2-methyl-3-aminopropionate (142.3 g; 0.65 mol) and ethyl acrylate (83.9 g; 0.94 mol) in ethanol (150 ml) was refluxed for 4 days. The solvent was removed in *vacuo*, and distillation of the residue gave 2b (164.8 g; 79 %), b.p. 165–167 °C/47 Pa. Anal. $\text{C}_{14}\text{H}_{19}\text{NO}_3$; C, H, N. IR (film): 3030 (w), 2980 (m), 2810 (m), 1730 (s), 1495 (w), 1455 (m) cm^{-1} . ^1H NMR (60 MHz, CCl_4): δ 7.18 (5 H, s), 4.2–3.7 (4 H, dq), 3.52 (2 H, s), 2.9–2.0 (7 H, m), 1.4–0.9 (9 H, dt+m).

(RS)-Ethyl N-methoxycarbonyl-N-(2-ethoxycarbonylethyl)-2-methyl-3-aminopropionate (3b). A solution of 2b (79.95 g; 250 mmol) and aqueous hydrochloric acid (62.5 ml; 4 M) in ethanol (200

ml) was hydrogenated (ca. 250 kPa) in a PARR hydrogenation apparatus by using a 10 % Pd-C catalyst (5.6 g). The reaction mixture was filtered and evaporated in *vacuo*. To an ice-cooled solution of the residue in water (100 ml) was added with stirring an iced solution of potassium carbonate (86.94 g; 630 mmol) in water (100 ml) followed by addition of methyl chloroformate (28.20 g; 300 mmol). Stirring was continued at 0 °C for 30 min and at 25 °C for 1 h. The mixture was extracted with ether (3 x 200 ml). The combined and dried (K_2CO_3) ether phases were evaporated in *vacuo* and the residue distilled to give 3b (46.4 g; 64 %), b.p. 143–145 °C/53 Pa. Anal. $\text{C}_{14}\text{H}_{21}\text{NO}_4$; C, H, N. IR (film): 2980 (m), 1730 (s), 1705 (s), 1435 (m), 1445 (m), 1410 (m) cm^{-1} . ^1H NMR (60 MHz, CCl_4): δ 4.02 (4 H, q, J 7 Hz), 3.59 (3 H, s) 3.5–3.1 (4 H, m), 3.0–2.0 (3 H, m), 1.4–0.9 (9 H, m).

(RS)-Ethyl 1-methoxycarbonyl-4-hydroxy-5-methyl-1,2,5,6-tetrahydropyridine-3-carboxylate (4b). To a suspension of sodium (1.84 g; 80 mmol) in benzene-xylene (10:1) (80 ml) was added ethanol (9.2 g; 200 mmol). When the sodium was dissolved a solution of 3b (23.2 g; 80 mmol) in benzene (40 ml) was added with stirring. The reaction mixture was left for 3 days at 25 °C. Upon addition of hydrochloric acid (25 ml; 4 M) the organic phase was dried (Na_2SO_4) and evaporated in *vacuo* to give crude 4b (14.3 g). CC [silica gel: 400 g; eluents: benzene containing ethyl acetate (5–15 %)] followed by distillation gave 4b (8.1 g; 42 %), b.p. 124–128 °C/40 Pa. Anal. $\text{C}_{14}\text{H}_{21}\text{NO}_4$; C, H, N. IR (film): 2980 (w), 2870 (w), 1710 (s), 1655 (m), 1620 (m), 1450 (m), 1410 (m) cm^{-1} . ^1H NMR (60 MHz, CCl_4): δ 12.17 (1 H, s), 4.6–3.8 (4 H, m), 3.8–3.0 (5 H, m), 2.7–2.1 (1 H, m), 1.6–0.7 (6 H, m).

(3RS,4SR)-Ethyl 1-methoxycarbonyl-4-hydroxypiperidine-3-carboxylate (5a). A solution of 4a¹⁵ (55.0 g; 0.24 mol) in ethanol (500 ml) was hydrogenated (ca. 10 MPa) by using a Ra-Ni W-2 catalyst (9 g). The filtered and evaporated reaction product was distilled to give 5a (50.0 g; 90 %), b.p. 145–149 °C/9 Pa, m.p. 51.5–53.5 °C. Anal. $\text{C}_{14}\text{H}_{21}\text{NO}_4$; C, H, N. IR (film): 3600–3200 (m), 2990–2850 (several bands, m), 1725 (s), 1705 (s) cm^{-1} .

(3RS,4SR)-Dimethyl 4-hydroxypiperidine-1,3-dicarboxylate (6a). A solution of 5a (26 g; 0.11 mol) in a methanolic solution of hydrogen chloride (500 ml; 5 %) was refluxed for 18 h. The evaporated reaction mixture was distilled to give 6a (21.0 g; 86 %), b.p. 143–146 °C/9 Pa, m.p. 68.5–70.5 °C. Anal. $\text{C}_{14}\text{H}_{21}\text{NO}_4$; C, H, N. IR (KBr): 3450 (s), 3000–2830 (several bands, m), 1720 (s), 1685 (s), 1670 (s) cm^{-1} .

(3RS,4SR)-3-Carboxy-4-hydroxypiperidinium chloride (7a). A solution of 6a (4.0 g; 18.4 mmol) in hydrochloric acid (40 ml; 5 M) was refluxed for 150 min. The evaporated oily reaction product was crystallized from water-glacial acetic acid [35 ml; 2.5] to give 7a (1.6 g;

determination analyses and MHz ^1H NMR described in a of compounds using sodium as an in-

48 %), m.p. 170.0–172.0 °C. Anal. C₉H₁₁ClNO₂: C, 51.1; H, 6.1; N, 10.1. IR (KBr): 3600–3350 (s), 3160–2740 (several bands, s), 2720–2400 (several bands, m), 1720 (s), 1600 (m) cm⁻¹. ¹H NMR (60 MHz, D₂O): δ 4.6–4.4 (1 H, m), 3.5–2.8 (3 H, m), 2.2–1.8 (2 H, m).

(3RS,4SR)-4-Hydroxypiperidine-3-carboxylic acid zwitterion (3a). To a solution of 7a (1.32 g; 1.0 mmol) in water (5 ml) was added a solution of triethylamine (1.06 g; 10.5 mmol) in ethanol (5 ml). To the filtered solution was added N,N-dimethylformamide (4 ml). Upon standing at 25 °C for 7 days 3a (890 mg; 61 %) was isolated, m.p. 253–255 °C (decomp.). Anal. C₉H₁₁NO₂: C, 51.1; H, 6.1. IR (KBr): 3210 (s), 3100–2200 (several bands, s), 1815 (s), 1550 (s), 1540 (s) cm⁻¹. pK_A values (H₂O, 22 °C): 3.42 ± 0.05; 10.02 ± 0.03.

(3RS,4SR,5SR)-Ethyl 1-methoxycarbonyl-4-hydroxy-5-methylpiperidine-3-carboxylate (5c) and (3RS,4SR,5RS)-ethyl 1-methoxycarbonyl-4-hydroxy-5-methylpiperidine-3-carboxylate (5d). A solution of 4b (4.0 g; 16 mmol) in ethanol (130 ml) was hydrogenated (ca. 3 MPa) by using an Ra-Ni W-2 catalyst (3 g) for 23 h. The filtered and evaporated reaction mixture was shown by TLC [eluent: benzene-ethyl acetate (4:1)] to consist of two compounds (*R*_F = 0.24 and 0.17). CC (silica gel: 200 g; eluents: benzene containing ethyl acetate (15–25 %)) of the crude product and ball-tube distillation of the separated components at 133 Pa (oven temperature 180 °C) gave pure 5c and 5d. 5c (1.45 g; 37 %) had m.p. 79.0–79.5 °C. Anal. C₁₁H₁₅NO₃: C, 51.1; H, 6.1. IR (KBr): 3470 (m), 2950 (m), 2900 (m), 1725 (s), 1685 (s), 1480 (s), 1455 (m), 1440 (m), 1415 (m) cm⁻¹. ¹H NMR (60 MHz, CDCl₃): δ 4.4–3.9 (m) and 3.67 (s) (a total of 7 H), 3.4–2.0 (5 H, m), 1.9–0.3 (7 H, m). 5d (1.12 g; 29 %) had m.p. 45–46 °C. Anal. C₁₁H₁₅NO₃: C, 51.1; H, 6.1. IR (film): 3470 (m), 2955 (m), 2925 (m), 1730 (s), 1700 (s), 1685 (s), 1480 (m), 1445 (m), 1415 (m) cm⁻¹. ¹H NMR (60 MHz, CDCl₃): δ 4.10 (q, *J* 7 Hz) and 3.9–2.2 (m) (a total of 12 H), 2.2–1.5 (1 H, m), 1.5–0.7 (6 H, m).

(3RS,4SR,5SR)-Dimethyl 4-hydroxy-5-methylpiperidine-1,3-dicarboxylate (6c). A solution of 5c (531 mg; 2.2 mmol) in methanolic hydrogen chloride (10 ml; 5 %) was refluxed for 17 h. Ball-tube distillation of the evaporated reaction product at 133 Pa (oven temperature 180 °C) gave 6c (472 mg; 93 %), m.p. 80.0–81.0 °C. The IR spectrum was almost identical with that of 5c.

(3RS,4SR,5SR)-3-Carboxy-4-hydroxy-5-methylpiperidinium chloride (7c). A solution of 6c (1.25 g; 5.4 mmol) in aqueous hydrochloric acid (50 ml; 5 M) was refluxed for 75 min. Evaporation of the reaction mixture *in vacuo* and recrystallization (water–acetic acid) of the residue gave 7c (486 mg; 46 %), m.p. 232.5–233.5 °C. Anal. C₉H₁₁ClNO₂: C, 51.1; H, 6.1. IR (KBr): 3400 (m), 3145 (m), 2940 (m), 2900 (m), 1720 (s), 1605 (m), 1465 (m), 1450 (m), 1420 (m) cm⁻¹. ¹H NMR (60 MHz, D₂O): δ 4.23

(1 H, slightly broadened s), 3.6–2.5 (5 H, m), 2.3–1.8 (1 H, m), 1.00 (3 H, d, *J* 7 Hz).

(3RS,4SR,5SR)-4-Hydroxy-5-methylpiperidine-3-carboxylic acid zwitterion (3c). To a solution of 7c (100 mg; 0.51 mmol) in water (1 ml) was added a solution of triethylamine (57 mg; 0.56 mmol) in ethanol (1 ml). The mixture was left at 4 °C for 17 h. Recrystallization (water–ethanol) of crude 3c gave pure 3c (46 mg; 51 %), m.p. 116.0–117.0 °C. Anal. C₉H₁₁NO₂: C, 51.1; H, 6.1. IR (KBr): 3700–3100 (s), 3100–2400 (several bands, m), 1590 (s), 1470 (m), 1395 (s) cm⁻¹. pK_A values (H₂O, 22 °C): 3.39 ± 0.02; 10.13 ± 0.04.

(3RS,4SR,5RS)-Dimethyl 4-hydroxy-5-methylpiperidine-1,3-dicarboxylate (6d). 6d was synthesized as described above for 5c by using 5d (471 mg; 1.9 mmol) as a starting material. Purification of the crude product by ball-tube distillation at 133 Pa (oven temperature 180 °C) gave 6d (404 mg; 92 %). The IR spectrum was almost identical with that of 5d.

(3RS,4SR,5RS)-3-Carboxy-4-hydroxy-5-methylpiperidinium chloride (7d). 7d was synthesized as described above for 7c by using 6d (1.25 g; 5.4 mmol) as a starting material. Recrystallization (water–acetic acid) of the crude product gave 7d (297 mg; 28 %), m.p. 211.5–212.0 °C. Anal. C₁₁H₁₅ClNO₃: C, 51.1; H, 6.1. IR (KBr): 3445 (m), 3110 (m), 3000 (m), 2945 (m), 1720 (s), 1585 (m), 1470 (m), 1435 (m) cm⁻¹. ¹H NMR (60 MHz, D₂O): δ 4.0–3.7 (1 H, m), 3.7–2.6 (5 H, m), 2.4–1.8 (1 H, m), 1.07 (3 H, d, *J* 7 Hz).

(3RS,4SR,5RS)-4-Hydroxy-5-methylpiperidine-3-carboxylic acid zwitterion (3d). 3d was synthesized as described above for 3c by using 7d (100 mg; 0.51 mmol) as a starting material. Recrystallization (water–ethanol) of the crude product gave 3d (28 mg; 31 %), m.p. 234.0–235.0 °C. Anal. C₉H₁₁NO₂: C, 51.1; H, 6.1. IR (KBr): 3600–3350 (m), 3350–2800 (several bands, s–m), 1730 (m), 1600 (s), 1470 (m), 1410 (s) cm⁻¹. pK_A values (H₂O, 22 °C): 3.29 ± 0.03; 9.99 ± 0.06.

Guvacine hydrobromide (3-carboxy-1,2,5,6-tetrahydropyridinium bromide) (9a). A solution of 6a (13.5 g; 62 mmol) in aqueous hydrobromic acid (60 ml; 48 %) was refluxed for 24 h. Upon cooling of the reaction mixture to 5 °C pure 9a (10.4 g; 30 %) crystallized, m.p. 290 °C (decomp.). Anal. C₉H₁₁BrNO₂: C, 51.1; Br, 31.1. IR (KBr): 3175 (m), 2950 (s), 2820 (m), 2620–2400 (several bands, w), 1720 (s), 1660 (m), 1580 (m) cm⁻¹. UV [methanol (log ε)]: 212 (4.00) nm. ¹H NMR (60 MHz, D₂O): δ 7.3–7.0 (1 H, m), 3.9–3.7 (2 H, q), 3.5–3.1 (2 H, t), 2.7–2.3 (2 H, m). pK_A values (H₂O, 22 °C): 3.50 ± 0.06; 9.85 ± 0.06.

(RS)-3-Carboxy-5-methyl-1,2,5,6-tetrahydropyridinium bromide (9b). A solution of 6c (517 mg; 1.99 mmol) in aqueous hydrobromic acid (1.5 ml; 48 %) was refluxed for 24 h. Upon cooling of the reaction mixture to 24 °C TLC-pure 9b crystallized (162 mg; 37 %) [*R*_F:

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37 %) [R_f:

1973) No. 5

0.22; eluent: 1-butanol – acetic acid – water (4:1:1)]. An analytical sample was recrystallized (water – acetic acid) to give ηb , m.p. 253.5 – 254.5 °C. Anal. C₁₁H₁₄BrNO₂; C, H, Br, N. IR (KBr): 3350 – 3300 (m), 3250 – 2500 (several bands, s – m), 1710 (s), 1660 (m), 1605 (m), 1400 (m) cm⁻¹. UV [methanol (log ϵ)]: 214 (4.00) nm. ¹H NMR (60 MHz, D₂O): δ 7.2 – 6.9 (1 H, m), 3.9 – 3.7 (2 H, broadened s), 3.7 – 3.2 (1 H, m), 3.1 – 2.5 (2 H, m), 1.17 (3 H, d, J 7 Hz).

(RS)- β -Methyl-1,2,5,6-tetrahydropyridine-3-carboxylic acid zwitterion (10). ηb was synthesized as described above for δc by using ηb (100 mg; 0.56 mmol) and a solution of triethylamine (63 mg; 0.62 mmol) in ethanol (2 ml). Recrystallization (water – ethanol) of the crude product gave ηb (37 mg; 47 %), m.p. 249.0 – 249.5 °C. Anal. C₁₁H₁₄NO₂; C, H, N. IR (KBr): 3600 – 3250 (m), 3200 – 2000 (several bands, m), 1660 (s), 1630 (s), 1550 (s), 1470 (m) cm⁻¹. UV [methanol (log ϵ)]: 210 (3.88) nm. pK_A values (H₂O, 23 °C): 3.29 ± 0.03; 9.85 ± 0.03.

(3RS,4SR)-Dimethyl 4-hydroxyperhydroazepine-1,3-dicarboxylate (12) and (3RS,4RS)-dimethyl 4-hydroxyperhydroazepine-1,3-dicarboxylate (13). A mixture of ηb and ηc was prepared from ηb (2.58 g; 10.5 mmol) in ethanol (15 ml) by using 1.5 g of catalyst as described above for $\delta c, d$, followed by reflux of a solution of the crude reduction product in methanolic hydrogen chloride (150 ml; 15 %) for 24 h. CC [silica gel: 180 g; eluents: benzene containing ethyl acetate (15 – 25 %)] of the evaporated reaction product followed by ball-tube distillation at 60 Pa (oven temperature 180 °C) of the isolated components gave ηb (1.63 g; 69 %), m.p. 55.5 – 57.5 °C. Anal. C₁₀H₁₇NO₃; C, H, N. IR (KBr): 3450 (m), 2950 (m), 2930 (m), 1735 (s), 1705 – 1675 (s), 1485 – 1440 (m), 1410 (m) cm⁻¹. 349 mg (14 %) of ηc was obtained. Anal. C₁₀H₁₇NO₃; C, H, N. IR (film): 3450 (m), 2950 (m), 2930 (m), 1735 (s), 1705 – 1680 (s), 1485 – 1440 (m), 1410 (m) cm⁻¹.

3-Carboxy-2,5,6,7-tetrahydro-1H-azepinium bromide (14). A solution of ηb (370 mg; 1.6 mmol) in hydrobromic acid (1 ml; 48 %) was reduced for 24 h. Evaporation in *vacuo* and recrystallization (water – isopropanol) of the residue gave ηb (225 mg; 63 %), m.p. 224.5 – 225.5 °C. Anal. C₁₀H₁₄BrNO₂; C, H, Br, N. IR (KBr): 3430 (m), 3150 – 2700 (several bands, s – m), 1715 (s), 1650 (m), 1575 (m), 1460 (m), 1420 (m) cm⁻¹. UV [methanol (log ϵ)]: 210 (4.00) nm. ¹H NMR (60 MHz, D₂O): δ 7.55 (1 H, t, J 7 Hz), 4.12 (2 H, s), 3.6 – 3.3 (2 H, m), 2.8 – 2.3 (2 H, m), 2.2 – 1.6 (2 H, m). pK_A values (H₂O, 23 °C): 3.71 ± 0.05; 10.20 ± 0.03.

(RS)-3-Carboxyperhydroazepinium bromide (15). A solution of ηb (490 mg; 2.2 mmol) in aqueous ethanol (150 ml; 35 %) was hydrogenated (ca. 300 kPa) in a PARR hydrogenation apparatus by using a 10 % Pd – C catalyst

(100 mg). Evaporation in *vacuo* and recrystallization of the residue (water – acetic acid) gave ηb (368 mg; 74 %), m.p. 145.0 – 146.5 °C. Anal. C₁₀H₁₄BrNO₂; C, H, Br, N. IR (KBr): 3700 – 3300 (m), 3250 – 2900 (s), 2860 (m), 2750 – 2400 (several bands, m – w), 1730 (s), 1580 (m) cm⁻¹. ¹H NMR (60 MHz, D₂O): δ 3.7 – 2.9 (5 H, m), 2.4 – 1.8 (6 H, m). pK_A values (H₂O, 23 °C): 3.33 ± 0.04; 10.58 ± 0.04.

(1R,2R,3S,5S)-(-)-Methyl 3-ethoxycarbonyl-

oxy-3-ethoxycarbonylnortropane-3-carboxylate (17). A solution of ecgonine hydrate (16) (3.0 g; 14.8 mmol) in methanolic hydrogen chloride (200 ml; 5 %) was refluxed for 4 h. Upon evaporation in *vacuo* an aqueous solution (7 ml) of the residue, adjusted to pH 10 by sodium hydroxide, was extracted with chloroform (3 × 30 ml). The combined, dried (Na₂CO₃), and evaporated organic phases were submitted to ball-tube distillation at 130 Pa (oven temperature 150 °C) to give ecgonine methyl ester (2.5 g; 85 %) as a TLC-pure oil [R_f: 0.11; eluent: 1-butanol – acetic acid – water (4:1:1)]. A solution of ecgonine methyl ester (2.5 g; 12.6 mmol) and ethyl chloroformate (14.0 g; 130 mmol) in 1,2-dichloroethane (70 ml) was refluxed for 3 days. Evaporation in *vacuo* and CC [silica gel: 180 g; eluents: dichloromethane containing ethyl acetate (20 – 50 %) and methanol (3 %)] of the oily residue followed by ball-tube distillation at 100 Pa (oven temperature 170 °C) gave ηb (1.63 g; 39 %), m.p. 92.0 – 93.5 °C. Anal. C₁₅H₂₂NO₂; C, H, N. [α]_D²⁵ – 48° (c 0.91, ethanol). IR (KBr): 3600 – 3100 (m), 2980 (m), 2890 (w), 1735 (s), 1710 (s), 1480 (m), 1435 (s) cm⁻¹. ¹H NMR (60 MHz, CCl₄): δ 5.2 – 4.7 (1 H, m), 4.7 – 4.3 (2 H, m), 4.3 – 3.8 (4 H, dq, each J 7 Hz), 3.66 (3 H, s), 3.1 – 2.8 (2 H, q), 2.7 – 2.2 (1 H, m), 2.0 – 1.8 (4 H, m), 1.5 – 1.2 (6 H, dt, each J 7 Hz).

(1R,5S)-(-)-2-Nortropene-2-carboxylic acid hydrobromide (18). A solution of ηb (835 mg; 2.5 mmol) in aqueous hydrobromic acid (6 ml; 48 %) was refluxed for 5 h. Evaporation in *vacuo* and recrystallization (2-propanol – ether) of the residue gave ηb (389 mg; 66 %), m.p. 273.5 – 280.5 °C. Anal. C₁₀H₁₄BrNO₂; C, H, Br, N. [α]_D²⁵ – 61° (c 0.98, water). IR (KBr): 3080 (s), 2990 (s), 2730 – 2440 (several bands, m – w), 1690 (s), 1640 (m), 1600 (m) cm⁻¹. UV [methanol (log ϵ)]: 212 (3.95) nm. ¹H NMR (60 MHz, D₂O): δ 7.2 – 7.0 (1 H, t, J 4 Hz), 4.8 – 4.6 (1 H, m), 4.5 – 4.2 (1 H, m), 3.4 – 2.4 (2 H, m), 2.4 – 1.9 (4 H, m). pK_A values (H₂O, 23 °C): 3.33 ± 0.03; 10.23 ± 0.05.

(1R,2R,5R)-(+)-Nortropane-2-carboxylic acid hydrobromide (19). ηb was prepared as described above for ηb from ηb (250 mg; 1.1 mmol) by using 50 mg of the catalyst. 203 mg (81 %) of ηb was obtained, m.p. 237.5 – 238.5 °C. Anal. C₁₀H₁₄BrNO₂; C, H, Br, N. [α]_D²⁵ + 12° (c 0.98, water). IR (KBr): 3600 – 3300 (m), 3120 (m), 3000 – 2830 (s), 2900 – 2400 (several bands, m – w), 1730 (s), 1610 (m) cm⁻¹.

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A computer-generated and "zoomed in" portion of synthetic DNA. It is displayed on an Evans and Sutherland PS340 via a "CPK" modeling algorithm implemented by Paul Isaacson. The coordinates are taken from file 172 on the Brookhaven Protein Data Bank listing, and the structure was determined by H. Drew and R. Dickerson.

FRONTISPICE

A computer-generated representation of serylglutamine, a dipeptide. Courtesy Evans and Sutherland.

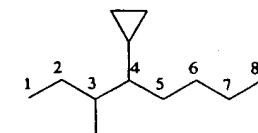
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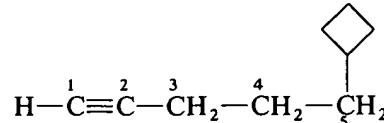
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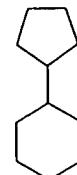
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4-cyclopropyl-3-methyloctane



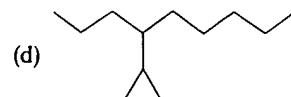
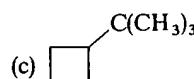
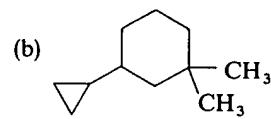
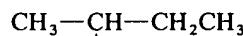
5-cyclobutyl-1-pentyne



cyclopentylcyclohexane

PROBLEM 3-14

Give a IUPAC name for each of the following compounds.



PROBLEM 3-15

Draw the structure and give the molecular formula for each of the following compounds.

(a) cyclododecane

(b) propylcyclohexane

(c) cyclopropyl cyclopentane

(d) 3-ethyl-1,1-dimethylcyclohexane

3-6D GEOMETRIC ISOMERISM IN CYCLOALKANES

Open-chain alkanes undergo rotations about their carbon-carbon single bonds, and they are free to assume any of an infinite number of conformations. Alkenes have rigid double bonds that prevent rotation, giving rise to *cis* and *trans* isomers with different orientations of the groups on the double bond. Cycloalkanes are similar to alkenes in this respect. A cycloalkane has two distinct faces. If two substituents point toward the same face, they are *cis*. If they point toward opposite faces, they are *trans*. These geometric isomers cannot interconvert without breaking and re-forming bonds.

Figure 3-15 compares the geometric isomers of 2-butene with those of 1,2-dimethylcyclopentane. You should make models of these compounds to convince

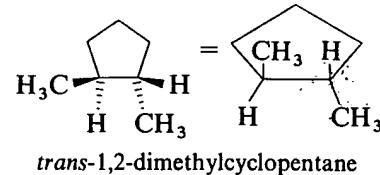
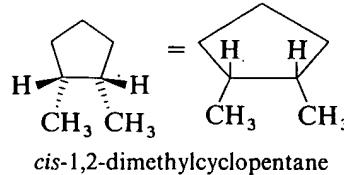
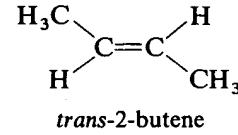
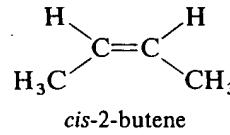


FIGURE 3-15 Like alkenes, cycloalkane rings are restricted from free rotation. Two substituents on a cycloalkane must be either on the same side (*cis*) or on opposite sides (*trans*) of the ring.

Both of these conformations require one group to be axial while the other is equatorial. The ethyl group is bulkier than the methyl group, so the conformation with the ethyl group equatorial is more stable. These chair conformations are in equilibrium at room temperature, and the one with the equatorial ethyl group predominates.

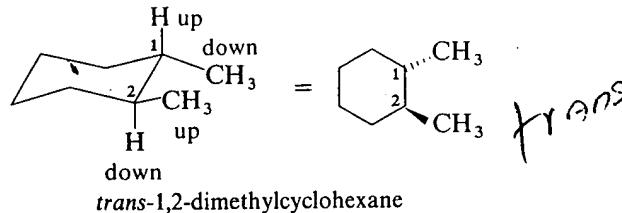
PROBLEM 3-26

Draw the two chair conformations of each of the following substituted cyclohexanes. In each case, label the more stable conformation.

- (a) *cis*-1-ethyl-2-methylcyclohexane
- (b) *trans*-1-ethyl-2-methylcyclohexane
- (c) *cis*-1-ethyl-4-isopropylcyclohexane
- (d) *trans*-1-ethyl-4-methylcyclohexane

Recognizing *cis* and *trans* isomers

Some students find it difficult to look at a chair conformation and tell whether a disubstituted cyclohexane is the *cis* isomer or the *trans* isomer. In the following drawing the two methyl groups appear to be oriented in similar directions. They are actually *trans* but are sometimes mistakenly identified as *cis*.



This ambiguity is eliminated by recognizing that each of the ring carbon atoms has two available bonds, one directed upward and one directed downward. In this drawing the methyl group on C1 is on the downward bond, and the methyl on C2 is on the upward bond. Because one is on a downward bond and one on an upward bond, their relationship is *trans*. A *cis* relationship would require both groups to be upward or both to be downward.

PROBLEM 3-27

Name each of the following compounds, remembering that two up bonds are *cis*; two down bonds are *cis*; one up bond and one down bond are *trans*.

